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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : C07D 243/12, 243/18, 243/24, 255/04, 257/10, 259/00, 487/00, 491/00, 495/00, A61K 31/55		A1	(11) International Publication Number: WO 98/00405 (43) International Publication Date: 8 January 1998 (08.01.98)						
(21) International Application Number: PCT/US97/11131 (22) International Filing Date: 25 June 1997 (25.06.97) (30) Priority Data: <table><tr><td>60/020,747</td><td>28 June 1996 (28.06.96)</td><td>US</td></tr><tr><td>9617894.2</td><td>28 August 1996 (28.08.96)</td><td>GB</td></tr></table> (71) Applicant (for all designated States except US): MERCK & CO., INC. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): LYNCH, Joseph, J., Jr. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). SALATA, Joseph, J. [US/US]; 126 East Lincoln Avenue, Rahway, NJ 07065 (US). (74) Common Representative: MERCK & CO., INC.; 126 East Lincoln Avenue, Rahway, NJ 07065 (US).		60/020,747	28 June 1996 (28.06.96)	US	9617894.2	28 August 1996 (28.08.96)	GB	(81) Designated States: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, HU, IL, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, US, UZ, VN, YU, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>	
60/020,747	28 June 1996 (28.06.96)	US							
9617894.2	28 August 1996 (28.08.96)	GB							
(54) Title: PHARMACEUTICAL PREPARATION									
(57) Abstract <p>A method of preventing, treating, terminating and protecting against cardiac arrhythmias, such as atrial, supraventricular and ventricular ectopy, tachycardia, flutter or fibrillation, including atrial, supraventricular and ventricular arrhythmias resulting from myocardial ischemic injury in a patient in need thereof, comprising administration of a selective I_{Ks} antagonist and a beta-adrenergic receptor blocking agent, administered in combined therapy either simultaneously, separately or sequentially is presented. Additionally, a pharmaceutical preparation comprising a selective I_{Ks} antagonist and a beta-adrenergic receptor blocking agent, wherein these compounds are administered simultaneously, separately or sequentially is presented.</p>									

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TITLE**PHARMACEUTICAL PREPARATION****BACKGROUND OF THE INVENTION**

The present invention relates to the co-administration, either simultaneously, separately or sequentially of a selective I_{K_S} antagonist and a beta-adrenergic receptor blocking agent for use in preventing, treating and terminating cardiac arrhythmias, such as atrial, supraventricular and ventricular ectopy, tachycardia, flutter or fibrillation, including atrial, supraventricular and ventricular arrhythmias resulting from myocardial ischemic injury in a patient in need thereof. This invention also relates to a pharmaceutical formulation which comprises a selective I_{K_S} antagonist and a beta-adrenergic receptor blocking agent along with a pharmaceutically acceptable carrier.

Arrhythmias often occur as complications to cardiac diseases such as myocardial infarction and heart failure. In a serious case, arrhythmias give rise to ventricular fibrillation and can cause sudden death.

Though various antiarrhythmic agents are now available on the market, agents exhibiting both satisfactory effects and high safety profiles, are not yet available for patients. For example, antiarrhythmic agents of Class I, according to the classification of Vaughan-Williams, which cause a selective inhibition of the maximum velocity of the upstroke of the action potential (V_{max}) are inadequate for preventing ventricular fibrillation. In addition, they have problems regarding safety, namely, they cause a depression of the myocardial contractility and have a tendency to induce arrhythmias due to an inhibition of the impulse conduction. Beta-adrenergic receptor blocking agent which belong to Class II are of limited value since their effects are either limited to a certain type of arrhythmia or are contraindicated because of their cardiac depressant properties in certain patients with cardiovascular disease. Their safety, however, is higher than that of the antiarrhythmic agents of Class I.

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Antiarrhythmic agents of Class III are drugs which cause a selective prolongation of the duration of the action potential without a significant depression of the Vmax. Until recently, drugs in this class were limited to sotalol and amiodarone, both of which have been shown to possess Class III properties. However, Sotalol also possesses Class II effects which may cause cardiac depression and be contraindicated in certain susceptible patients. Amiodarone is severely limited by side effects. Drugs of this class are expected to be effective in preventing ventricular fibrillations. Pure Class III agents, by definition, are not considered to cause myocardial depression or an induction of arrhythmias due to the inhibition of the action potential conduction as seen with Class I antiarrhythmic agents.

Recently, a novel group of Class III agents have been disclosed which antagonize the I_{Ks} channel found in heart muscle. These compounds I_{Ks} channel antagonists are effective in treating and preventing all types of arrhythmias including ventricular and atrial (supraventricular) arrhythmias. These novel compounds are disclosed and claimed in U.S. Patent Application, Serial Nos. 08/411,240; 08/516,467; and 08/516,226 which are hereby incorporated by reference. These novel compounds are especially useful for controlling reentrant arrhythmias and preventing sudden death due to ventricular fibrillation. These compounds are also effective in treating and preventing impaired cardiac pump functions.

In the treatment of arrhythmia, I_{Ks} antagonists have demonstrated effectiveness when delivered orally in amounts ranging from about 0.01 to about 1 mg per kg of body weight per day, in a single dose or in 2 to 4 divided doses.

The activity of the compounds described herein as anti-arrhythmic agents is measured by their ability to block the I_{Ks} and I_{Kr} currents as determined by the following test protocol.

Outward potassium currents are measured in single guinea pig ventricular myocytes using a whole-cell voltage clamp technique described in detail elsewhere (Sanguinetti and Jurkiewicz, 1990, Two components of cardiac delayed rectifier K^+ current: differential

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sensitivity to block by Class III antiarrhythmic agents. J. Gen Physiol. 96: 195-215). Myocytes are isolated by enzymatic (collagenase and protease) digestion of Langendorff perfused hearts. Single cells are then voltage clamped using 1 mm square-bore pipettes filled with 0.5 M Kgluconate, 25 mM KCl, 5 mM K(2)ATP. Cells are bathed in a solution containing, in mM: 132 NaCl, 4KCl, 1.2 MgCl₂, 10 HEPES, 10 glucose: pH 7.2, temp. 35°C.

Each cell is maintained at a holding potential of -50 mV. Test depolarizations are applied as voltage ramps from -85 to -50 mV, and as steps to -10 mV (0.5 s) and +50 mV (1.0 s). I_{K1} is measured as peak outward current during the voltage ramp. I_{Kr} is measured as tail currents upon repolarization from -10 mV to -50 mV. I_{Ks} is measured as time-dependent current during the pulse to +50 mV. Currents are measured during control, then after exposure to drug at two different concentrations.

Employing this test the compounds described herein as selective I_{Ks} channel antagonists, have an IC₅₀ of less than 100 nM as I_{Ks} antagonists. The compounds of this invention are at least 10 times more potent in the blockade of I_{Ks} than of blockade of I_{Kr} .

Beta-adrenergic receptor blocking agents, or "beta-blockers", are a class of pharmaceutically active compounds which decrease the positive chronotropic, positive inotropic, bronchodilator and vasodilator responses caused by beta-adrenergic receptor agonists. The magnitude of this decreased response is proportional to the existing sympathetic tone and the concentration of beta-blocker at the receptor sites. Beta-adrenergic receptor blockage is said to reduce cardiac output in both healthy subjects and patients with heart disease. While the mechanism of antihypertension effects of beta-adrenergic receptor blocking agents has not been established, possible mechanisms of action include reduction in cardiac output, reduction in plasma renin activity, and central nervous system sympatholytic action. The administration of beta-adrenergic receptor antagonists has been shown effective in reducing the incidence of mortality and sudden death in postinfarction

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patients (Yusaf et al., Prog Cardiovasc Dis 17: 335-371, 1985; Lau et al., N Eng J Med 327: 248-254, 1992).

While both selective I_{K_S} channel blockers and beta-adrenergic receptor blocking agents have been proven effective when administered separately, it is considered to be in the best interest of the patient to reduce the amount of these compounds provided to the patient. Any reduction of one or the other compound would be considered helpful, but this is particularly true of beta-adrenergic receptor blocking agents which are known to have significant side effects in some humans.

SUMMARY OF THE INVENTION

A method is presented for use in preventing, treating and terminating cardiac arrhythmias, such as atrial, supraventricular and ventricular ectopy, tachycardia, flutter or fibrillation, including atrial, supraventricular and ventricular arrhythmias resulting from myocardial ischemic injury in a patient in need thereof which comprises the co-administration, either simultaneously, separately or sequentially of a selective I_{K_S} antagonist and a beta-adrenergic receptor blocking agent. This invention also relates to a pharmaceutical formulation which comprises a selective I_{K_S} antagonist and a beta-adrenergic receptor blocking agent along with a pharmaceutically acceptable carrier.

DETAILED DESCRIPTION OF THE INVENTION

A method is presented for use in preventing, treating and terminating cardiac arrhythmias, such as atrial, supraventricular and ventricular ectopy, tachycardia, flutter or fibrillation, including atrial, supraventricular and ventricular arrhythmias resulting from myocardial ischemic injury in a patient in need thereof which comprises the co-administration, either simultaneously, separately or sequentially of a selective I_{K_S} antagonist and a beta-adrenergic receptor blocking agent. This invention also relates to a pharmaceutical formulation which comprises a selective I_{K_S} antagonist and a beta-adrenergic receptor blocking agent along with a pharmaceutically acceptable carrier.

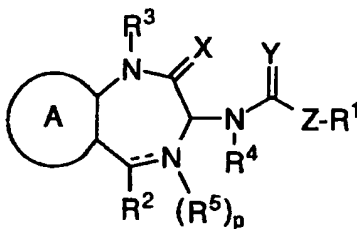
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By a "selective I_{K_S} antagonist" is meant those compounds which when studied in the test disclosed above have an IC_{50} of less than 100 nM as I_{K_S} blockers. The compounds of this invention are at least 10 times more potent in the blockade of I_{K_S} than of blockade of I_{K_R} .

Beta-adrenergic receptor blocking agents are compounds which decrease the positive chronotropic, positive inotropic, bronchodilator and vasodilator responses caused by beta-adrenergic receptor agonists. The magnitude of this decreased response is proportional to the existing sympathetic tone and the concentration of beta-adrenergic receptor blocking agent which reaches the receptor sites.

Examples of compounds which fit the definition of beta-adrenergic receptor blocking agent include but are not limited to timolol, sotalol, esmolol, cateolol, propranolol, betaxolol, penbutolol, metoprolol, acebutolol, atenolol, metoprolol, pindolol, and bisoprolol, and their salts, hydrates, solvates and any crystal forms in which they may occur.

Examples of compounds which fit the definition of selective I_{K_S} antagonists include, but are not limited to, the following:



I

or a pharmaceutically acceptable salt thereof, wherein

- A is
- 1) thieno,
 - 2) pyrido, or
 - 3) benzo either unsubstituted or substituted with $-NH_2$, $-NHSO_2$ (C_{1-3} alkyl), C_{1-3} alkyl or C_{1-3} alkoxy;

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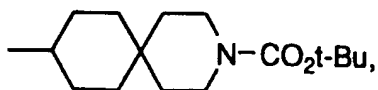
- X is
- 1) =O,
 - 2) =S,
 - 3) =N-NH₂,
 - 4) =N-OH or
 - 5) =H₂;

- Y is
- 1) =O,
 - 2) =N-CN or
 - 3) =H₂;

- Z is
- 1) C₁-6 alkylene, either straight or branch chain and either unsubstituted or substituted with phenyl or spiro-piperidine,
 - 2) C₂-4 alkenylene, either straight or branch chain,
 - 3) -(CH₂)_m-W-(CH₂)_n- wherein m and n are independently 0, 1, 2, 3 or 4 and W is -O-, -S- or -NH,
 - 4) 4-(5-methylisoxazole-3-yl),
 - 5) C₃-6 cycloalkylene, or
 - 6) single bond;

p is 0 or 1;

- R¹ is
- 1) phenyl, either unsubstituted or substituted with one or two substituents selected from
 - a) -NO₂,
 - b) -Cl, Br, F, or I,
 - c) -CF₃,
 - d) -C₁-3 alkyl,
 - e) -C₁-3 alkoxy,
 - f) -CN,
 - g) -methylenedioxy,
 - 2) C₅-7 cycloalkyl,
 - 3)



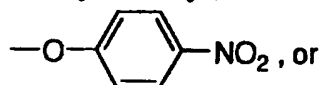
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- 4) mono- or bicyclic heterocyclyl of 5 to 10 members one or two of which are sulfur, nitrogen or oxygen, the remaining being carbon, such as 2-thienyl, 2-furanyl, 2-indolyl, 2-quinoxoliny, or 2-(2,3-dihydro benzofuranyl)
- 5) C₁₋₃ alkyl, or
- 6) indan-5-yl;

- R² is
- 1) phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,
 - 2) C₁₋₆ alkyl, either straight or branched chain, and either unsubstituted or substituted with C₁₋₃ alkoxy or C₁₋₃ alkoxy-C₁₋₃ alkoxy,
 - 3) C₅₋₇ cycloalkyl,
 - 4) 2- or 3-furyl,
 - 5) 1-methylpiperidin-2-yl, or
 - 6) if R² is phenyl, the 2-position of the phenyl can be joined to the 4-position nitrogen of the diazepine ring through a carbonyl group and the double bond between the 4-nitrogen and the 5-carbon becomes a single bond;

- R³ is
- 1) hydrogen or
 - 2) C₁₋₃ alkyl either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃, or
 - 3) -CF₃;

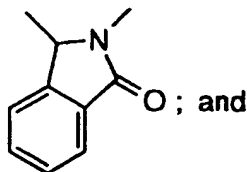
- R⁴ is
- 1) hydrogen,
 - 2) C₁₋₆ alkyl, the chain of carbon atoms of which can be interrupted by one or two non-adjacent oxygen atoms and which is either unsubstituted or substituted with C₁₋₃ alkoxycarbonyl, -OH or



- 3) tetrazol-5-yl;

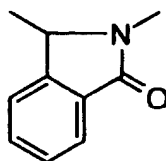
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R⁵ is hydrogen or oxygen or is joined to R² to form the partial structure:



the bond represented by ----- is:

- 1) a double bond when p is zero or when p is 1 and R⁵ is oxygen, or
- 2) a single bond when R⁵ is hydrogen or R⁵ is joined to R² to form the partial structure:



This invention is meant to include the individual diastereomers where such exist and mixtures thereof and enantiomers and mixtures of the enantiomers.

The pharmaceutically acceptable salts of the compounds of Formulas I include the conventional non-toxic salts or the quarternary ammonium salts of the compounds of Formula I formed, e.g., from non-toxic inorganic or organic acids. For example, such conventional non-toxic salts include those derived from inorganic acids such as hydrochloric, hydrobromic, sulfuric, sulfamic, phosphoric, nitric and the like; and the salts prepared from organic acids such as acetic, propionic, succinic, glycolic, stearic, lactic, malic, tartaric, citric, ascorbic, pamoic, maleic, hydroxymaleic, phenylacetic, glutamic, benzoic, salicylic, sulfanilic, 2-acetoxybenzoic, fumaric, toluenesulfonic, methanesulfonic, ethane disulfonic, oxalic, isethionic, and the like.

The pharmaceutically acceptable salts of the present invention can be synthesized from the compounds of Formula I which

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contain a basic or acidic moiety by conventional chemical methods. Generally, the salts are prepared by reacting the free base or acid with stoichiometric amounts or with an excess of the desired salt-forming inorganic or organic acid or base in a suitable solvent or various combinations of solvents.

One embodiment of this invention are novel compounds useful in the novel method of treatment of this invention wherein:

A is benzo;

X and Y are oxygen;

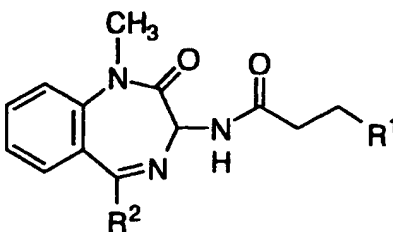
R³ is methyl;

R⁴ is hydrogen; and

R² is C₁-6 alkyl.

Specific novel compounds representative of this embodiment are those of the following structure and specified in Table I:

TABLE I

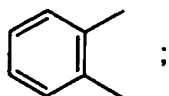


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<u>R¹</u>	<u>R²</u>
2,4-diClPh	-CH ₃
2,4-diClPh	-C ₂ H ₅
2,4-diClPh	-t-Bu
4-CF ₃ Ph	i-C ₃ H ₇
cyclohexyl	i-C ₃ H ₇
2,4-diClPh	i-C ₃ H ₇

Another embodiment of the compounds useful in the novel method of treatment of this invention is that wherein:

A is



X and Y are oxygen;

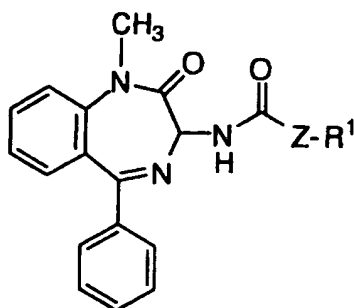
R³ is methyl;

R⁴ is hydrogen; and

R² is phenyl.

A class of novel compounds within this embodiment is that with structural formula:

- 11 -



wherein

Z is C₁-6 alkylene or a bond and

R¹ is phenyl, phenyl substituted with -Cl, -Br, -I, -F, or -CF₃, or R¹ is cyclohexyl.

Specific novel compounds representative of this class are those depicted in the following Table II:

TABLE II

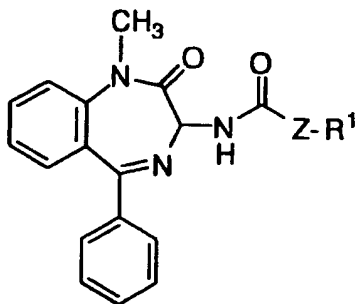
Z	R ¹
-(CH ₂) ₂ -	2,4-diClPh
-(CH ₂) ₂ -	4-ClPh
-(CH ₂) ₂ -	2,4-diFPh
-(CH ₂) ₂ -	2-ClPh
-(CH ₂) ₂ -	4-CF ₃ Ph
-CH ₂ -	4-CF ₃ Ph
-(CH ₂) ₂ -	3-CF ₃ Ph
-(CH ₂) ₂ -	2-CF ₃ Ph
-(CH ₂) ₂ -	cyclohexyl
-	cyclohexyl

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TABLE II (Cont'd)

<u>Z</u>	<u>R¹</u>
-(CH ₂) ₃ -	cyclohexyl
-CH ₂ -	cyclohexyl
-(CH ₂) ₂ -	Ph
-CH ₂ -	Ph
-(CH ₂) ₂ -	4-CNPh
-(CH ₂) ₂ -	3-ClPh
-(CH ₂) ₃ -	Ph
-(CH ₂) ₂ -	3-CNPh
-(CH ₂) ₃	2-thienyl

Another class of novel compounds within this embodiment is that with structural formula:



wherein Z is C₂-4 alkenylene and R¹ is phenyl or phenyl substituted with -Cl, -Br, -F, -I, -CF₃, C₁-3 alkyl, C₁-3 alkoxy or methylenedioxy.

Specific novel compounds representative of this class are those depicted in the following Table III:

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TABLE III

<u>Z</u>	<u>R¹</u>
-CH=CH-	4-NO ₂ Ph
-CH=CH-	2,4-diClPh
-CH=CH-	3-ClPh
-CH=CH-	2-ClPh
-CH=CH-	2,4-diFPh
-CH=CH-	2,6-diClPh
-CH=CH-	4-CF ₃ Ph
-CH=CH-	2-BrPh

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TABLE III (Cont'd)

<u>Z</u>	<u>R¹</u>
-CH=CH-	4-IPh
-CH=CH-	4-BrPh
$\begin{array}{c} \text{-C=CH-} \\ \\ \text{CH}_3 \end{array}$	Ph
-CH=CH-	Ph*
-CH=CH-	3,4-diClPh
-CH=CH-	4-CH ₃ Ph
-CH=CH-	4-CH ₃ OPh
-CH=CH-	3,4-methylenedioxyPh
-CH=CH-	3-BrPh

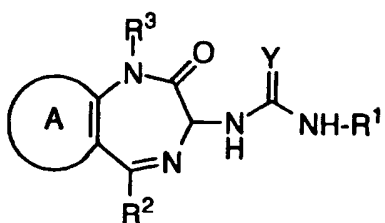
*This compound is disclosed in U.S. Patent 4,820,834

A third embodiment of the compounds useful in the novel method of treatment of this invention is that wherein:

Z is -NH-.

Compounds representative of this embodiment are those disclosed in the following Table IV.

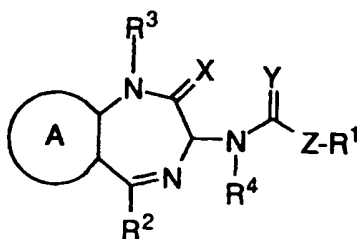
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TABLE IV

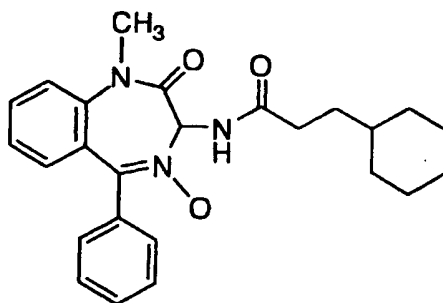
A	R ¹	R ²	R ³	Y
benzo	3-CH ₃ Ph	Ph		O
benzo	2,4-diClPh	Ph	-CH ₃	O
benzo	3-CH ₃ Ph		n-C ₃ H ₇	O
benzo	-CH ₂ Cyclohexyl	Ph	-CH ₃	=N-CN
benzo	3-CH ₃ Ph	Ph	-CH ₃	O
benzo	5-indanyl	Ph		O
	3-CH ₃ Ph	Ph	-CH ₃	O

Other specific compounds included within the broadest genus but not included in one of the embodiments previously described are as shown in Table V.

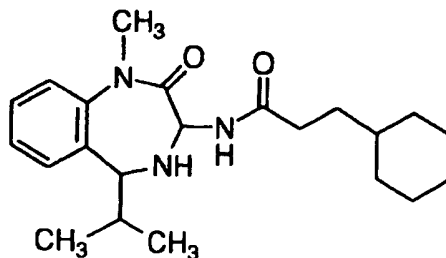
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TABLE V

Representative of compounds wherein p is 1 is the compound of structural formula:

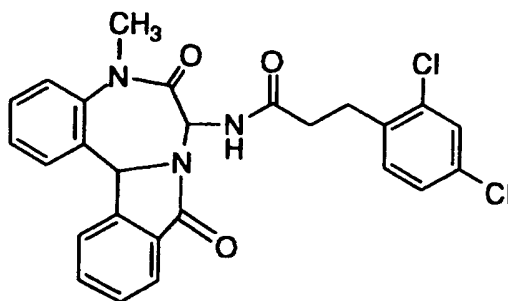


Representative of compounds wherein the bond between the 4 and 5 positions is a single bond is the compound of structural formula:

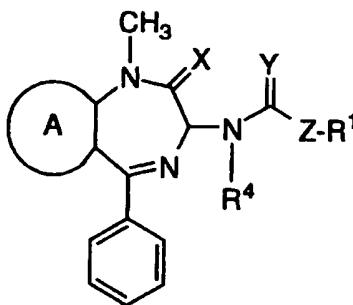


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Representative of compounds wherein the bond ----- represents a single bond and R^5 is joined to R^2 is the compound of structural formula:

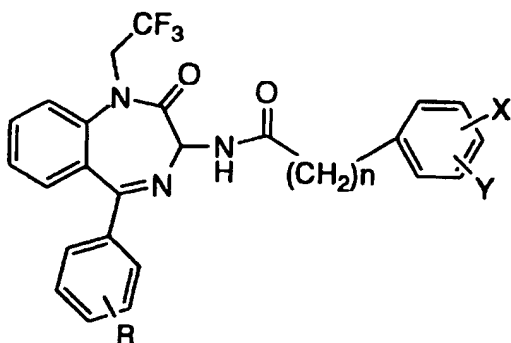


Another embodiment of this invention is a group of compounds, active in the novel method of treatment of this invention, which are novel compounds *per se*. These novel compounds are depicted in the following Table VI.



Another embodiment of this invention is a group of compounds which are active in the novel method of treatment of this invention. These compounds are depicted as follows:

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where

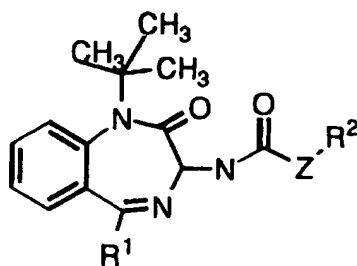
X and Y are independently hydrogen, chloro, fluoro, bromo, iodo, or trifluoromethyl and

n is 0, 1 or 2;

R is hydrogen, fluoro, chloro, bromo, iodo, or trifluoromethyl, methyl, or methoxy; and

the racemates, mixtures of enantiomers, individual diastereomers or individual enantiomers with all isomeric forms and pharmaceutically acceptable salts, hydrates or crystal forms thereof, which are antiarrhythmic agents.

Yet another embodiment of this invention is a group of compounds which are active in the novel method of treatment of this invention. These compounds are depicted as follows:



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R¹ and R² are independently

- 1) **phenyl, either unsubstituted or substituted with one or two substituents selected from**
 - a) **-NO₂, OH,**
 - b) **-Cl, Br, F, or I,**
 - c) **-CF₃,**
 - d) **-C₁₋₃ alkyl,**
 - e) **-C₁₋₃ alkoxy,**
 - f) **-CN,**
 - g) **-methylenedioxy, and**

Z is

- 1) **C₁₋₆ alkyl, either straight or branched chain ,**
- 2) **substituted C₁₋₆ alkyl, either straight or branched chain, wherein the substituents are selected from F, OH, NO₂,**
- 3) **C₂₋₄ alkenylene, either straight or branched chain,**
- 4) **-(CH₂)_m-W-(CH₂)_n- wherein m and n are independently 0, 1, 2, 3 or 4 and W is -O-, -S- or -NH,**
- 5) **C₃₋₆ cycloalkane,**
- 6) **C₃₋₆ cycloalkylene, or**
- 7) **single bond;**

The selective I_{Ks} blockers of the present invention have the pharmacological properties required for antiarrhythmic agents of Class III, namely they demonstrate prolongation of QTc-interval , and dose dependent increases in ventricular refractoriness. This is accomplished without effecting heart rate, mean arterial pressure and PR and QRS intervals. Modest increases in LV+dP/dt (left ventricular change in pressure with time) is observed. Further, these compounds suppress the induction of PVS (Programmed Ventricular Stimulation) induced ventricular tachyarrhythmias.

Individually, these compounds are effective in treating

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and preventing all types of arrhythmias including ventricular, atrial and supraventricular arrhythmias. The compounds of the present invention are especially useful for controlling reentrant arrhythmias and prevent sudden death due to ventricular fibrillation. These compounds are also effective in treating and preventing impaired cardiac pump functions.

In the novel method of this invention of treating arrhythmia, a selective I_{Ks} antagonist is administered in an amount ranging from about .0001 to about 10 mg per kg of body weight per day, preferably from about .0001 to about 2 mg per kg of body weight per day, and more preferably, when intravenous delivery of the compounds is employed, from about 0.0003 to about 0.3 mg per kg of body weight per day, or when given orally from about 0.01 to about 1 mg per kg of body weight per day, in a single dose or in 2 to 4 divided doses of each compound. The beta-adrenergic receptor blocking agent is administered in an amount ranging from about 1 mg per day to about 300 mg per day and more preferably from about 2 mg/day to about 250 mg per day.

The activity of the compounds described herein as antiarrhythmic agents is measured by their ability to block the I_{Ks} and I_{Kr} currents as determined by the following test protocol.

Outward potassium currents are measured in single guinea pig ventricular myocytes using a whole-cell voltage clamp technique described in detail elsewhere (Sanguinetti and Jurkiewicz, 1990, Two components of cardiac delayed rectifier K^+ current: differential sensitivity to block by Class III antiarrhythmic agents. J. Gen Physiol. 96: 195-215). Myocytes are isolated by enzymatic (collagenase and protease) digestion of Langandorf perfused hearts. Single cells are then voltage clamped using 1 mm square-bore pipettes filled with 0.5 M Kgluconate, 25 mM KCl, 5 mM K(2)ATP. Cells are bathed in a solution containing, in mM: 132 NaCl, 4KCl, 1.2 $MgCl_2$, 10 HEPES, 10 glucose: pH 7.2, temp. 35°C.

Each cell is maintained at a holding potential of -50 mV. Test depolarizations are applied as voltage ramps from -85 to -50 mV, and as steps to -10 mV (0.5 s) and +50 mV (1.0 s). I_{K1} is measured as

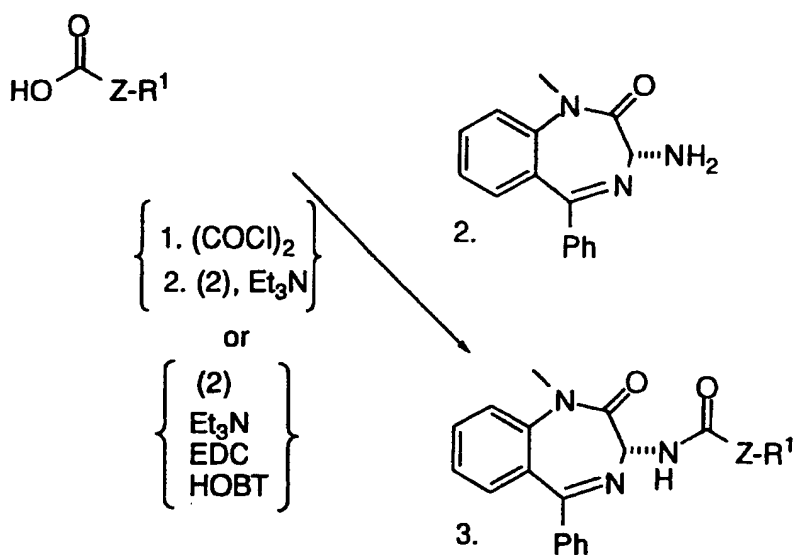
- 21 -

peak outward current during the voltage ramp. I_{K_r} is measured as tail currents upon repolarization from -10 mV to -50 mV. I_{K_s} is measured as time-dependent current during the pulse to +50 mV. Currents are measured during control, then after exposure to drug at two different concentrations.

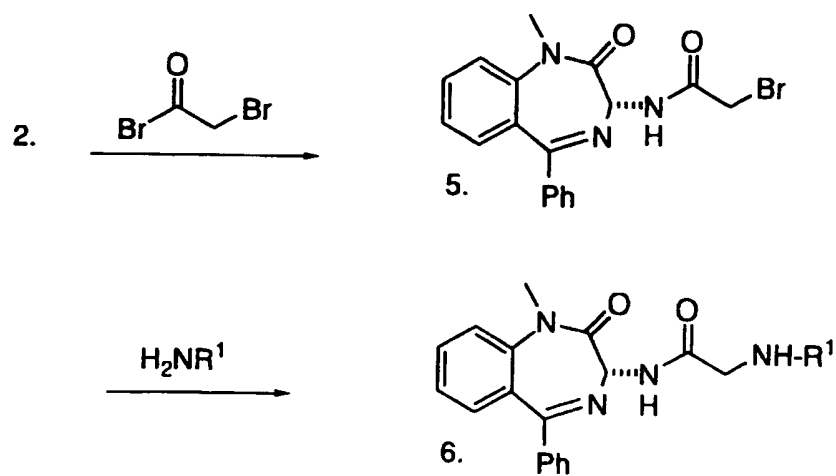
Employing this test the compounds described herein as selective I_{K_s} blockers have an IC_{50} of less than 100 nM as I_{K_s} blockers. The compounds of this invention are at least 10 times more potent in the blockade of I_{K_s} than of blockade of I_{K_r} .

Typical synthetic schemes employed in making the compounds herein are illustrated below.

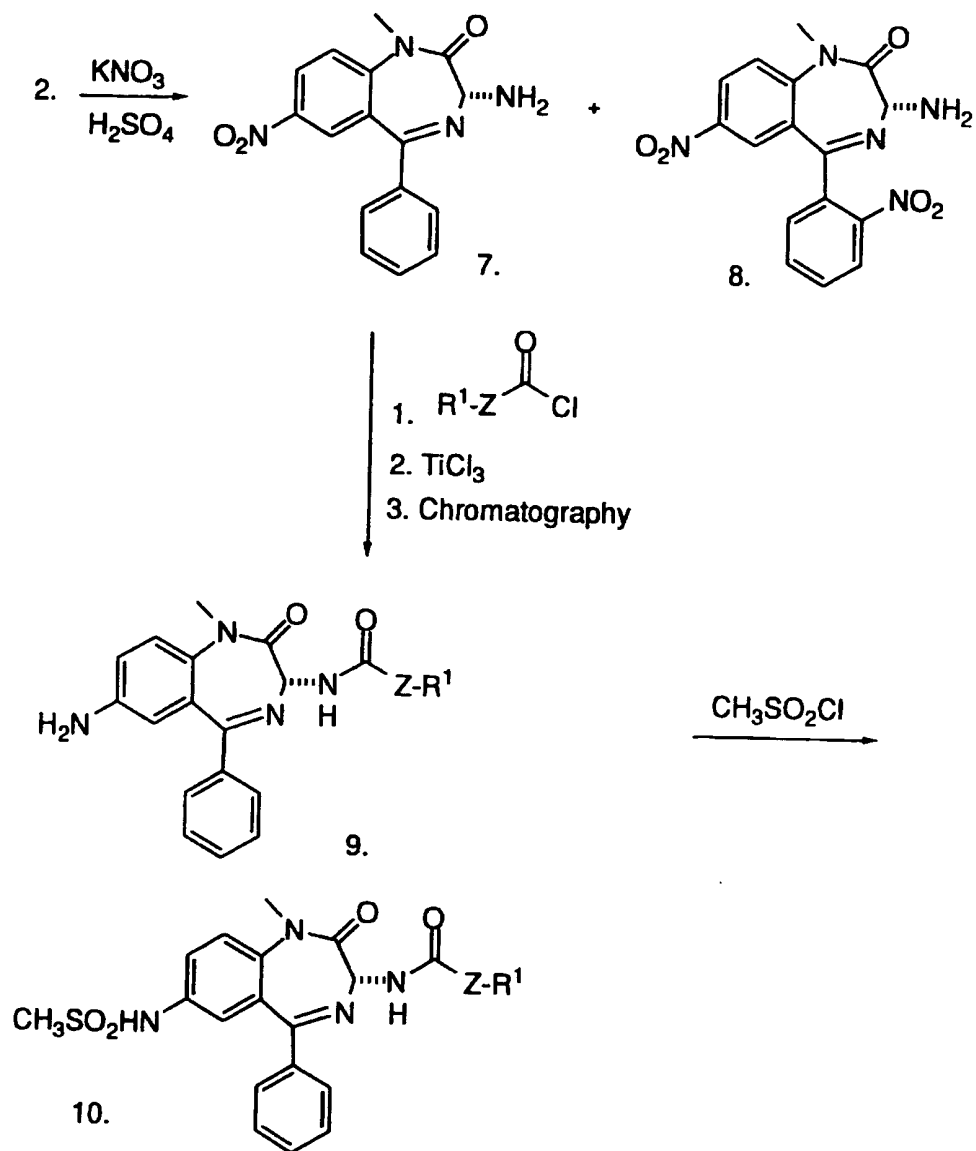
SCHEME 1



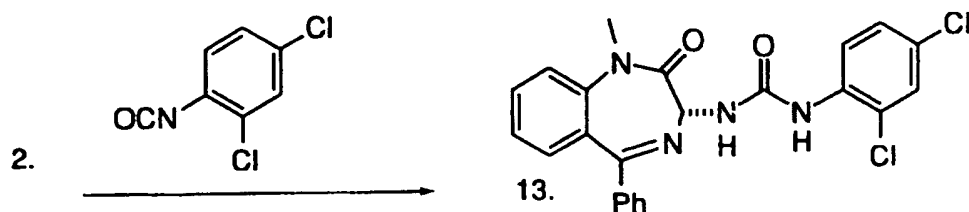
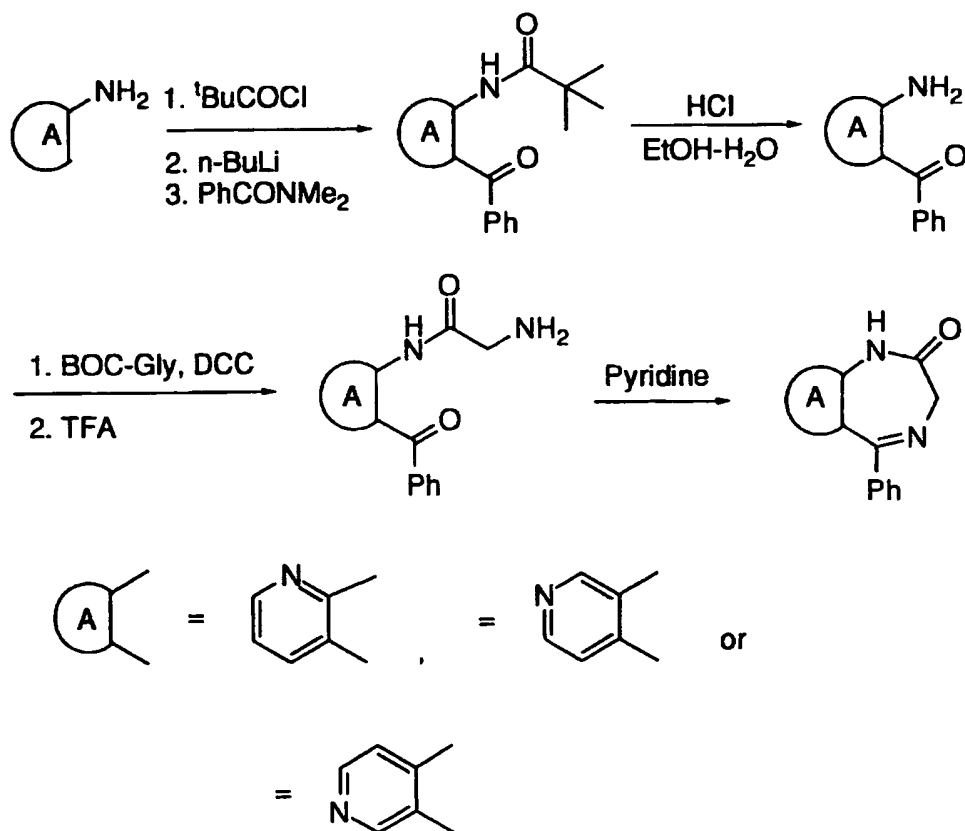
- 22 -

SCHEME 2

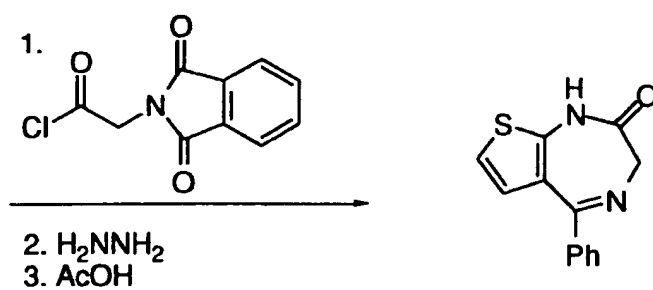
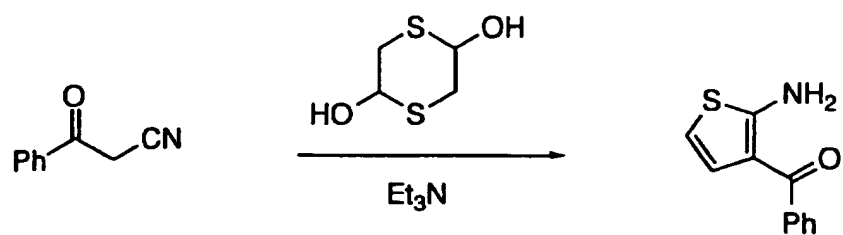
- 23 -

SCHEME 3

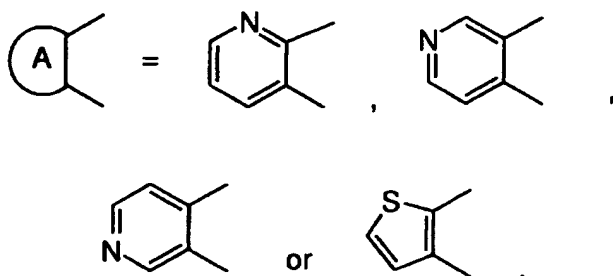
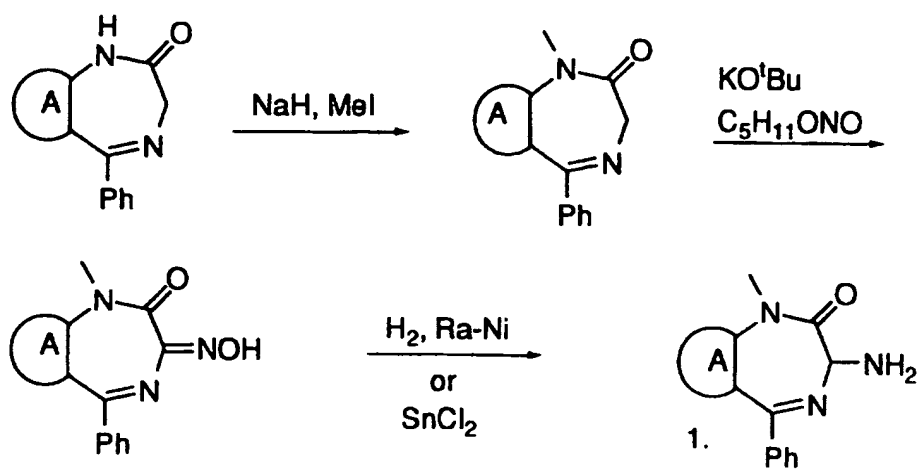
- 24 -

SCHEME 4SCHEME 5

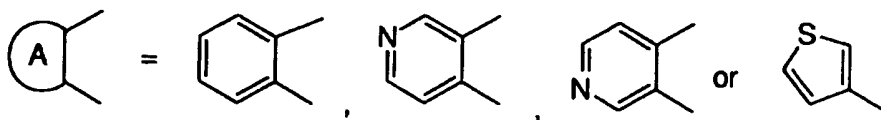
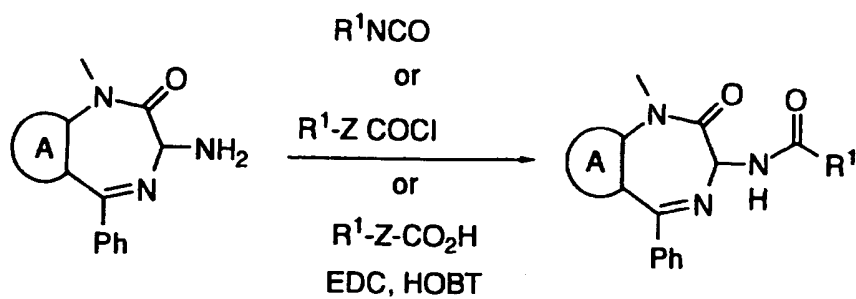
- 25 -

SCHEME 6

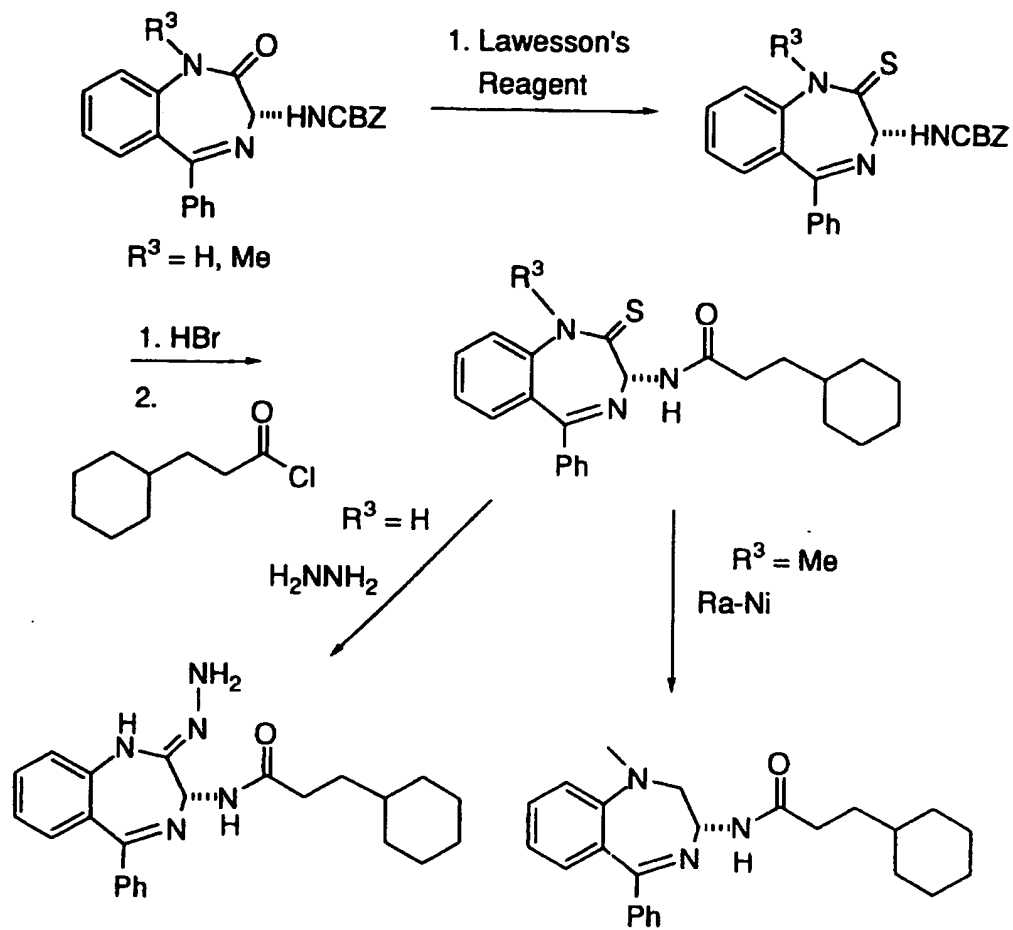
- 26 -

SCHEME 7

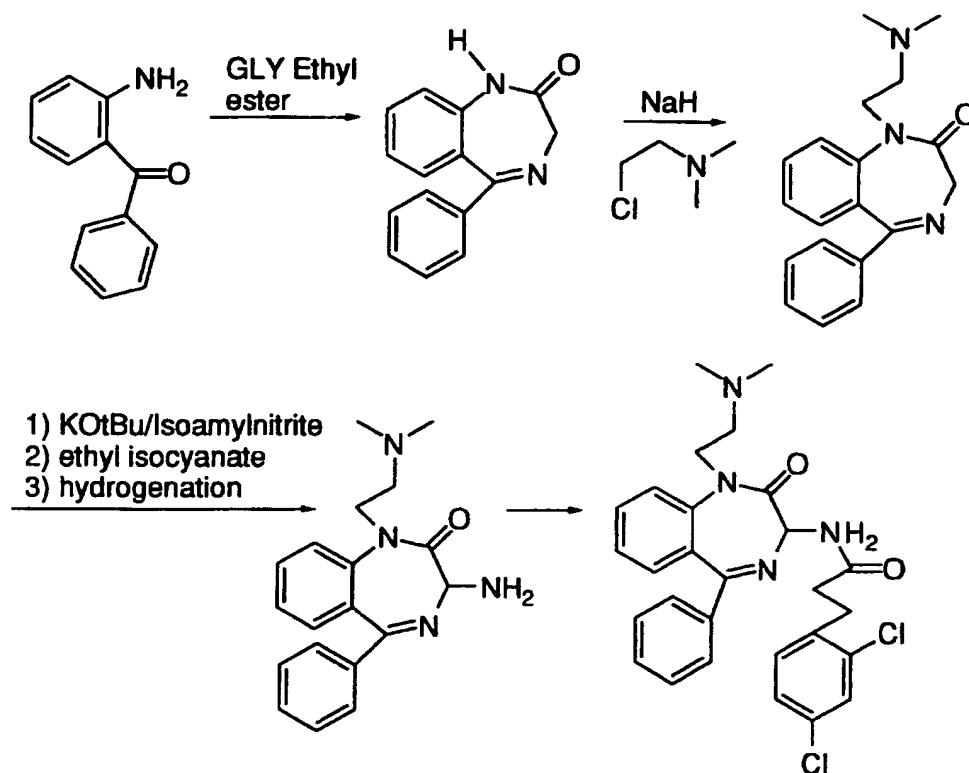
- 27 -

SCHEME 8

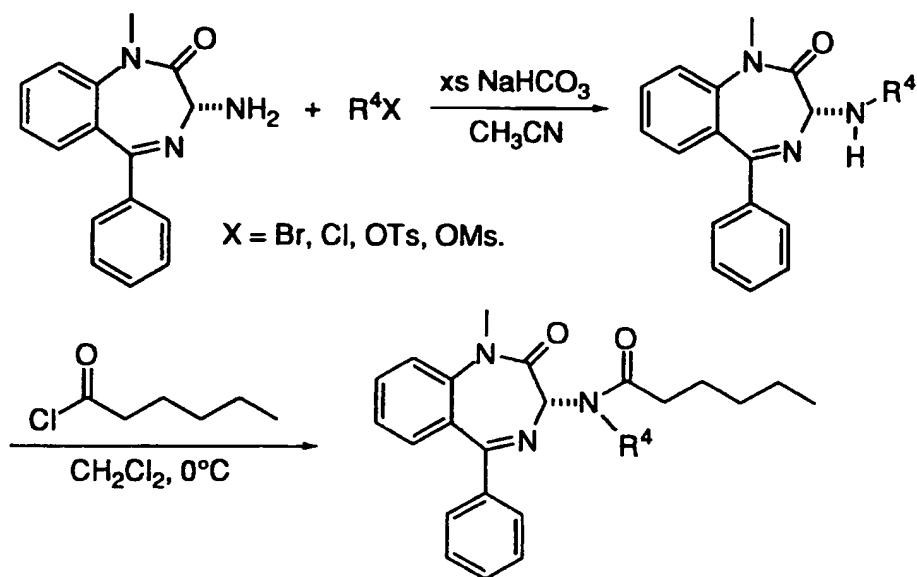
- 28 -

SCHEME 9

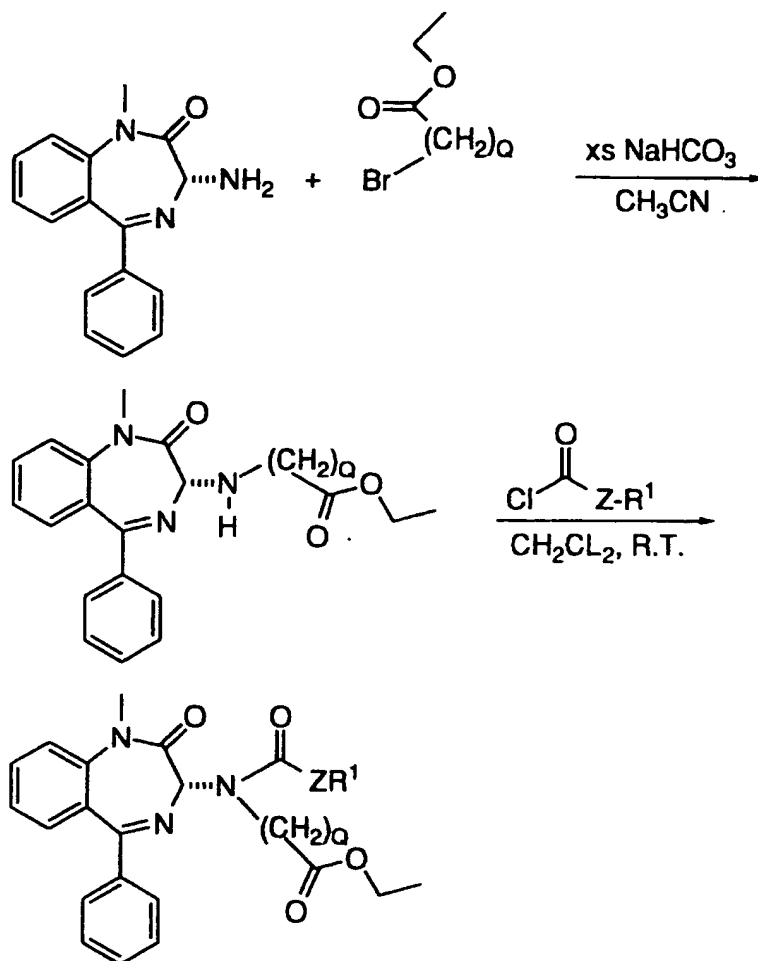
- 29 -

SCHEME 10

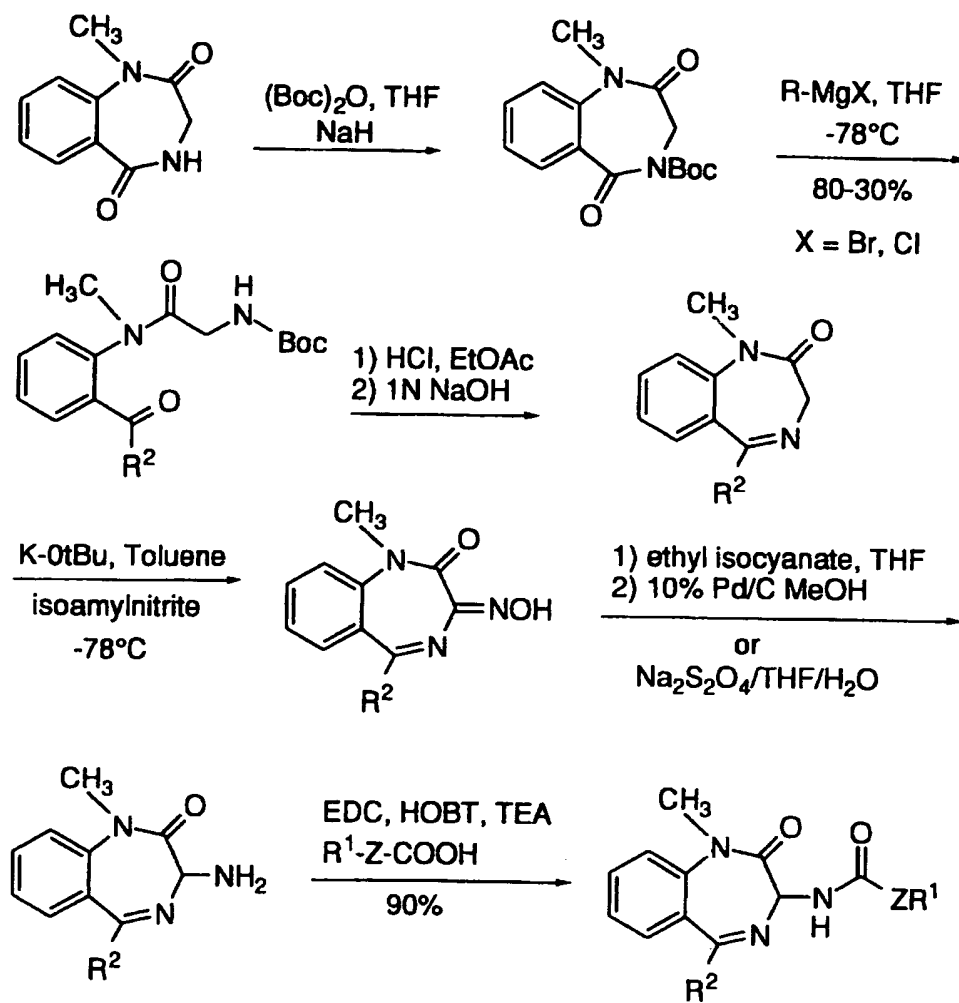
- 30 -

SCHEME 11

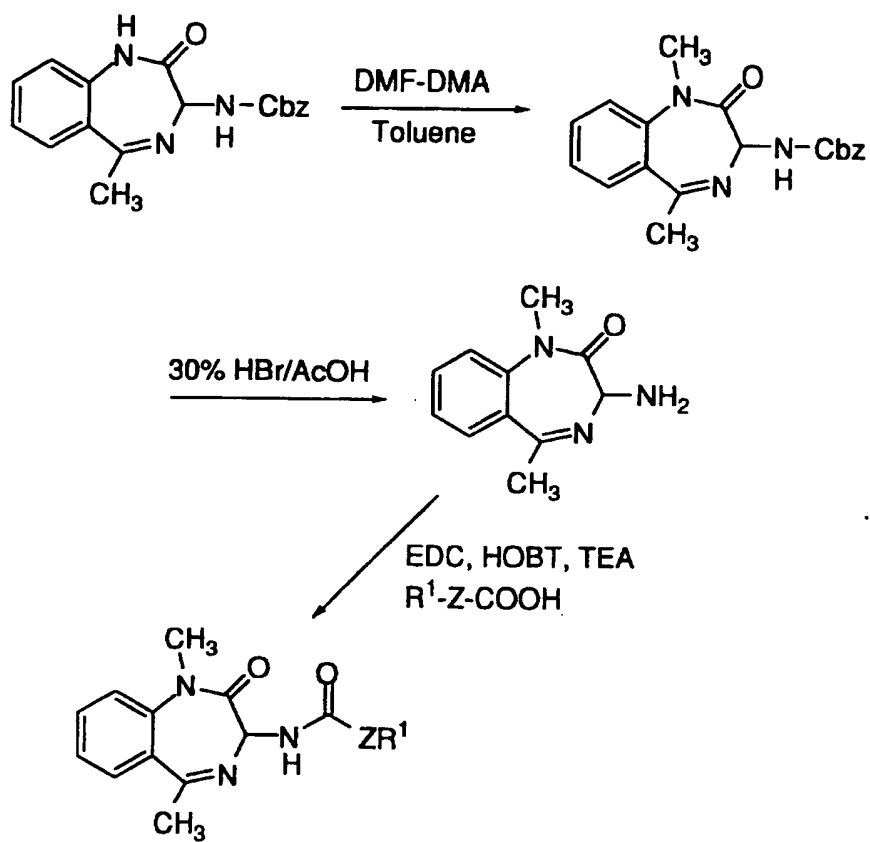
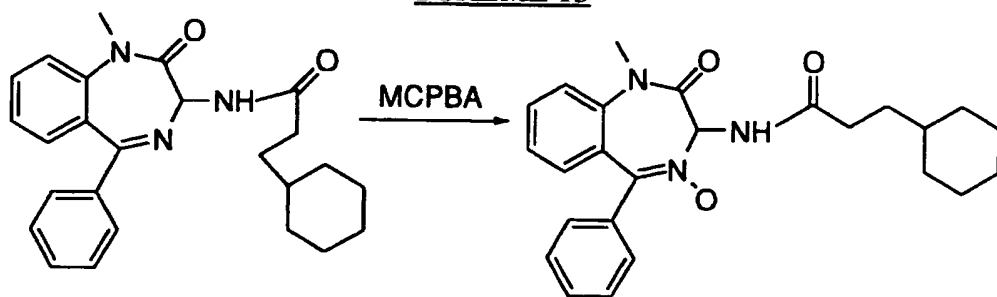
- 31 -

SCHEME 12

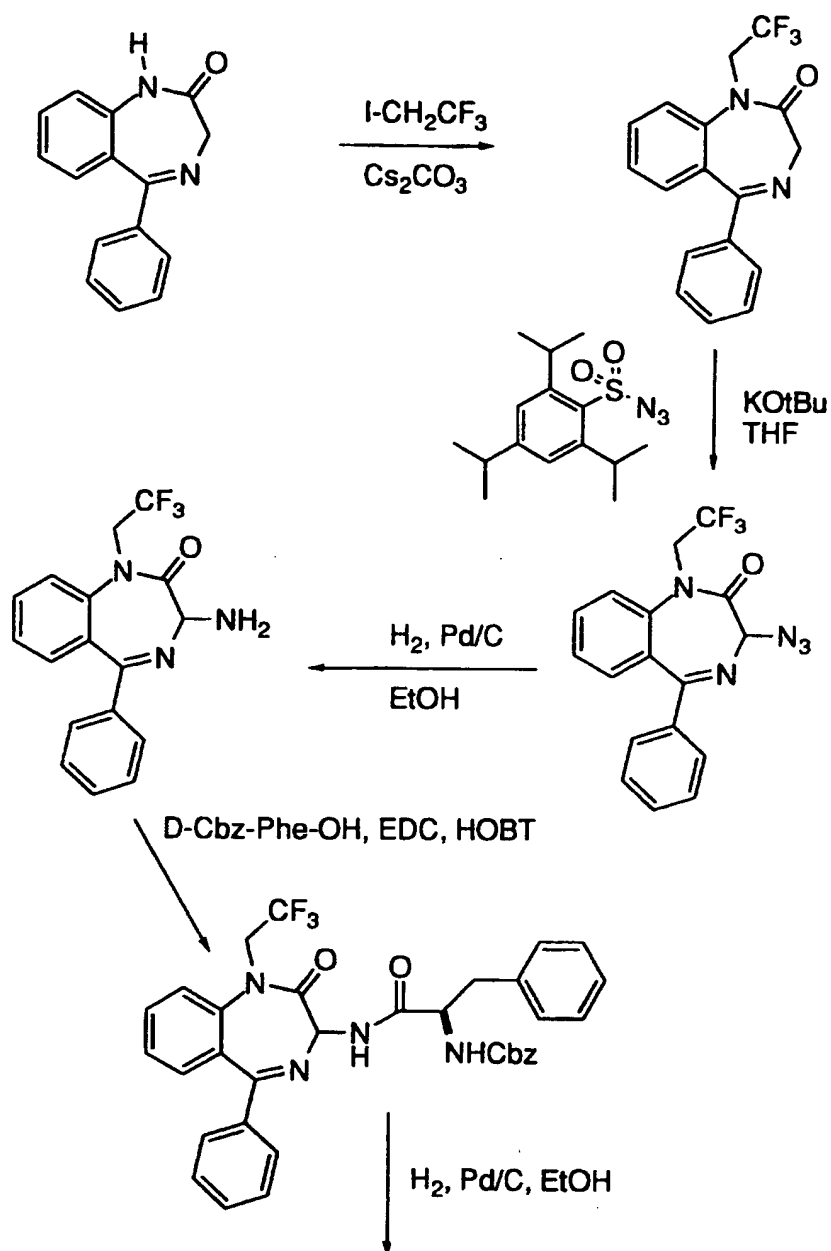
- 32 -

SCHEME 13

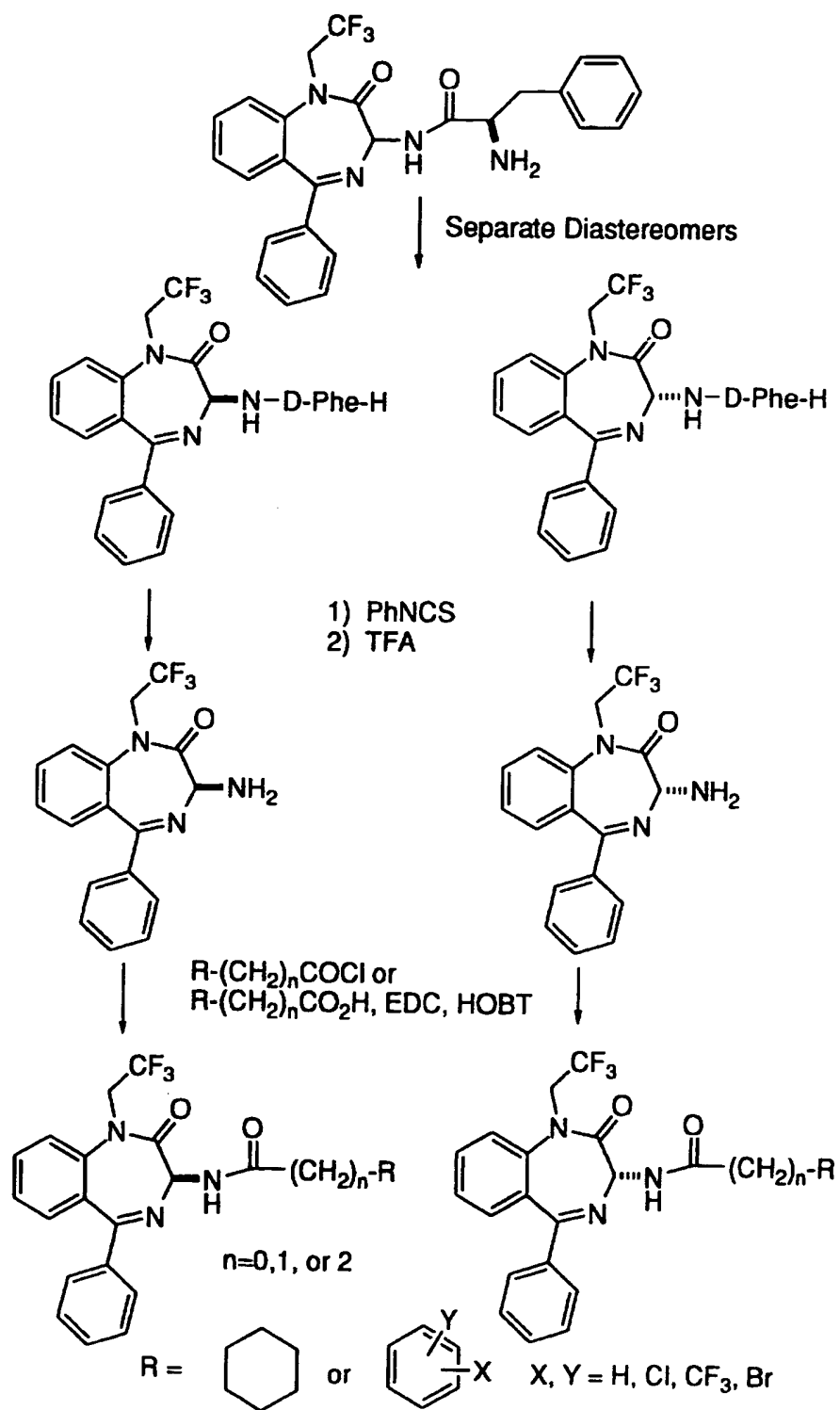
- 33 -

SCHEME 14SCHEME 15

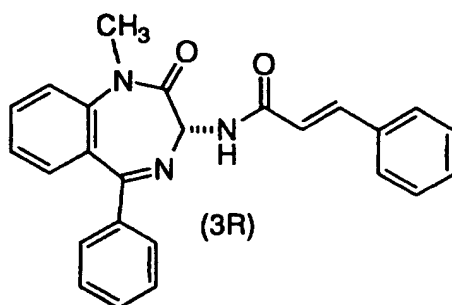
- 34 -

SCHEME 16

- 35 -



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EXAMPLESEXAMPLE 1(E)-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-phenyl-2-propenamide

A solution of (E)-3-phenyl-2-propenoyl chloride (367 mg, 2.2 mmol) in methylene chloride (1 mL) was added to a solution of 3(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (*J. Org. Chem.* 1987, 52, 3232-3239) (531 mg, 2.0 mmol) and triethylamine (307 mL, 225 mg, 2.2 mmol) in methylene chloride (10 mL). The mixture was stirred at room temperature for 25 min. and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with CH₂Cl₂/Et₂O (95:5) and the residue was triturated with Et₂O. The solid was collected and dried *in vacuo* at 70°C to give (E)-(+)-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-phenyl-2-propenamide as a colorless solid (170 mg, 21%), m.p. 140-142°C, [α]_D +86.7° (c=0.173, CH₂Cl₂).

¹H (CDCl₃) 7.70-7.26 (16H, m), 6.63 (1H, d, J 15.6 Hz), 5.68 (1H, d, J 8.3 Hz), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₂₁N₃O₂·0.15 (C₂H₅)₂O:

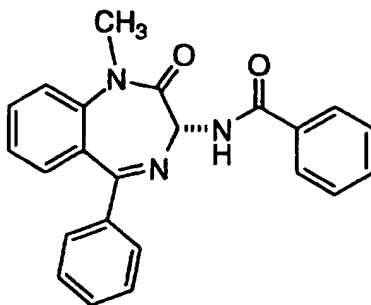
C, 75.63; H, 5.58; N, 10.33.

Found: C, 75.29; H, 5.57; N, 10.33%.

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Employing the procedure substantially as described above, but substituting an appropriate acid chloride for the (E)-3-phenyl-2-propenoyl chloride, the following compounds were prepared:

EXAMPLE 2



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]benzamide

m.p. 224-225°C, $[\alpha]_D^{+89.2^\circ}$ (c = 0.141, CH₂Cl₂).

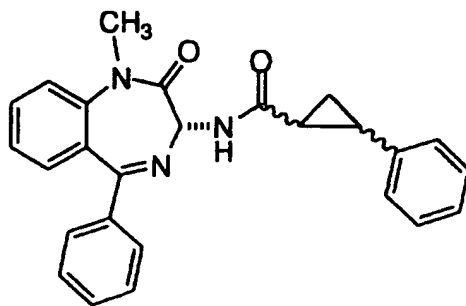
¹H (CDCl₃) 8.04 (1H, d, J 8.1 Hz), 7.96 (2H, d, J 6.8 Hz), 7.64-7.36 (10H, m), 7.27 (2H, t, J 7.6 Hz), 5.74 (1H, d, J 7.8 Hz), and 3.51 (3H, s).

Anal. Calcd. for C₂₃H₁₉N₃O₂·0.20H₂O:

C, 74.06; H, 5.24; N, 11.26.

Found: C, 74.13; H, 5.12; N, 11.16%.

EXAMPLE 3



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First diastereoisomer to elute:

(-)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzo-diazepin-3-yl](trans-2-phenyl-1-cyclopropane)carboxamide

m.p. 180-181°C, $[\alpha]_D -155.8^\circ$ (c = 0.434, CH₂Cl₂).

δ_H (CDCl₃) 7.62-7.09 (15H, m), 5.59 (1H, d, J 8.1 Hz), 3.47 (3H, s), 2.52-2.45 (1H, m), 1.90-1.84 (1H, m), 1.69-1.56 (1H, m), and 1.38-1.32 (1H, m).

Anal. Calcd. for C₂₆H₂₃N₃O₂·0.25H₂O:

C, 75.43; H, 5.72; N, 10.15.

Found: C, 75.38; H, 5.64; N, 9.94%.

Second diastereoisomer to elute:

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzo-diazepin-3-yl](trans-2-phenyl-1-cyclopropane)carboxamide

m.p. 104-107°C, $[\alpha]_D +328.2^\circ$ (c = 0.098, CH₂Cl₂).

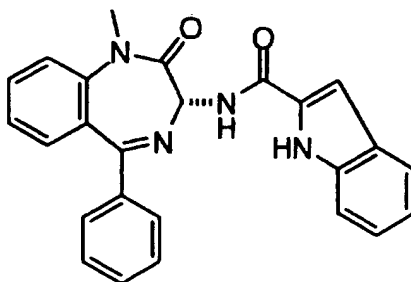
δ_H (CDCl₃) 7.62-7.13 (15H, m), 5.60 (1H, d, J 8.3 Hz), 3.48 (3H, s), 2.59-2.54 (1H, m), 1.93-1.87 (1H, m), 1.62-1.56 (1H, m, overlaps with water), and 1.33-1.25 (1H, m).

Anal. Calcd. for C₂₆H₂₃N₃O₂·0.50H₂O·0.45PhCH₃:

C, 76.13; H, 5.95; N, 9.14.

Found: C, 76.10; H, 5.94; N, 9.17%.

EXAMPLE 4



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(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-1H-indole-2-carboxamide

m.p. 167-177°C, $[\alpha]_D^{+113^\circ}$ (c = 1.103, CH₂Cl₂).

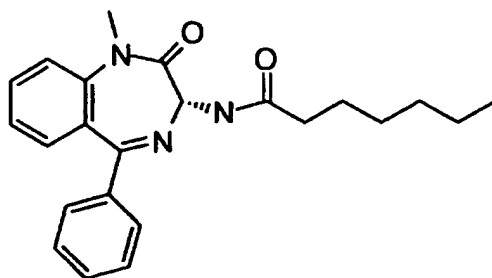
¹H (CDCl₃) 9.15 (1H, br s), 8.10 (1H, d, J 9.0 Hz), 7.75-7.10 (14H, m), 5.75 (1H, d, J 9.0 Hz), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₂₀N₄O₂:

C, 73.51; H, 4.94; N, 13.72.

Found: C, 73.31; H, 4.80; N, 13.62%.

EXAMPLE 5



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]heptanamide

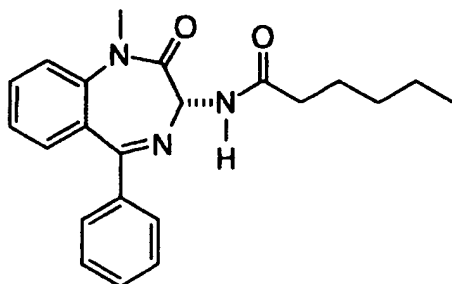
m.p. 49-54°C, $[\alpha]_D^{+69.5^\circ}$ (c=1.000, MeOH).

Anal. Calcd. for C₂₃H₂₇N₃O₂·0.40H₂O:

C, 71.81; H, 7.28; N, 10.92.

Found: C, 71.90; H, 7.09; N, 10.85%.

EXAMPLE 6



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(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]hexanamide

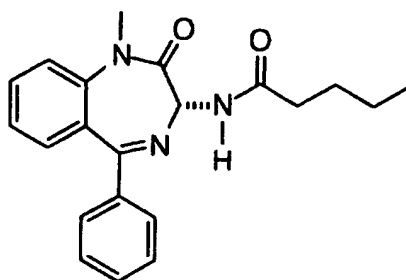
$[\alpha]_D^{+72.6^\circ}$ ($c=0.920$, MeOH).

Anal. Calcd. for $C_{22}H_{25}N_3O_2$:

C, 72.70; H, 6.93; N, 11.56.

Found: C, 72.44; H, 6.75; N, 11.25%.

EXAMPLE 7



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]pentanamide

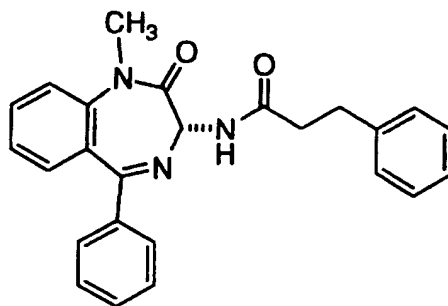
$[\alpha]_D^{+68.2^\circ}$ ($c=1.310$, MeOH).

Anal. Calcd. for $C_{21}H_{23}N_3O_2 \cdot 0.25CHCl_3$:

C, 68.21; H, 6.26; N, 11.26.

Found: C, 68.2; H, 6.29; N, 11.17%.

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EXAMPLE 8(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-phenylpropanamide

Oxalyl chloride (158 mL, 230 mg, 1.81 mmol) was added to a mixture of 3-phenylpropanoic acid (249 mg, 1.66 mmol) and DMF (1 drop) in THF (10 mL) and the mixture was stirred at room temperature for 40 min. 3(R)-Amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (*J. Org. Chem.* 1987, 52, 3232-3239) (400 mg, 1.51 mmol) and triethylamine (252 mL, 183 mg, 1.81 mmol) were added and the mixture was stirred at room temperature for 18 h. The mixture was poured into saturated aqueous sodium hydrogen carbonate (20 mL) and extracted with ethyl acetate (3 x 20 mL). The combined organic fractions were dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with CH₂Cl₂/Et₂O (95:5) and the residue was recrystallized from toluene/hexane to give (+)-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-phenylpropanamide as a colorless solid (380 mg, 63%), m.p. 179°C, [α]_D +100.4° (c = 0.225, CH₂Cl₂).

¹H (CDCl₃) 7.62-7.57 (2H, m), 7.47-7.21 (13H, m), 5.54 (1H, d, J 8.1 Hz), 3.47 (3H, s), 3.03 (2H, t, J 7.8 Hz), and 2.73-2.67 (2H, m).

Anal. Calcd. for C₂₅H₂₃N₃O₂·0.15H₂O:

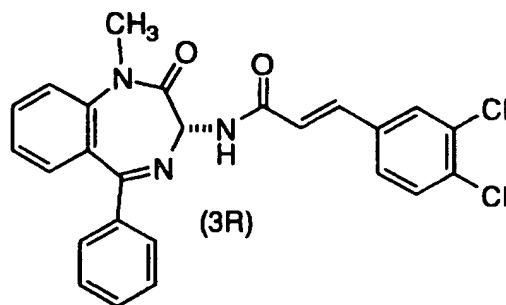
C, 75.04; H, 5.87; N, 10.50.

Found: C, 75.06; H, 5.78; N, 10.55%.

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Employing the procedure substantially as described above, but substituting an appropriate carboxylic acid for the 3-phenylpropanoic acid, the following compounds were prepared:

EXAMPLE 9



E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(3,4-dichlorophenyl)-2-propenamide

m.p. 145-147°C, $[\alpha]_D^{+77.8^\circ}$ (c=0.126, CH₂Cl₂).

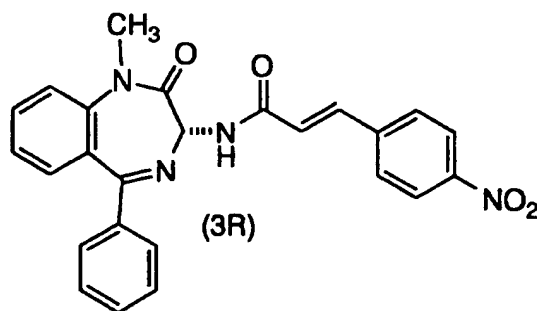
¹H (CDCl₃) 7.64-7.25 (14H, m), 6.61 (1H, d, J 15.6 Hz), 5.65 (1H, d, J 8.0 Hz), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₁₉N₃O₂Cl₂:

C, 64.67; H, 4.12; N, 9.05.

Found: C, 64.57; H, 4.25; N, 9.01%.

EXAMPLE 10



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E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzo-diazepin-3-yl]3-(4-nitrophenyl)-2-propenamide

m.p. 165-166 °C, $[\alpha]_D^{25} +80.5^\circ$ ($c=0.126$, CH_2Cl_2).

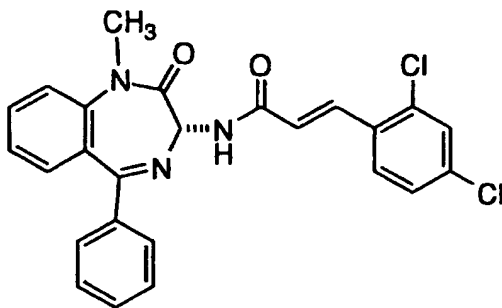
^1H (CDCl₃) 8.26 (1H, d, J 8.8 Hz), 7.74-7.28 (13H, m), 6.76 (1H, d, J 15.6 Hz), 5.66 (1H, d, J 8.0 Hz), and 3.51 (3H, s).

Anal. Calcd. for C₂₅H₁₉N₄O₄:

C, 68.17; H, 4.58; N, 12.72.

Found: C, 68.25; H, 4.65; N, 12.57%.

EXAMPLE 11



E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]3-(2,4-dichlorophenyl)-2-propenamide

m.p. 137-139°C, $[\alpha]_D^{25} +66.0^\circ$ ($c=0.144$, CH_2Cl_2).

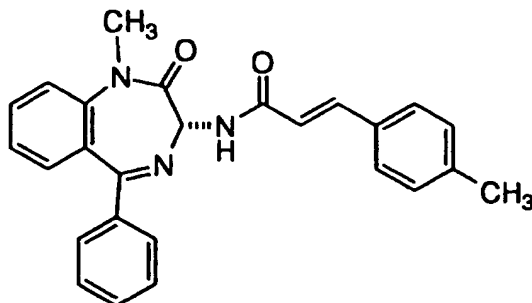
^1H (CDCl₃) 8.02 (1H, d, J 15.6 Hz), 7.73-7.26 (13H, m), 6.66 (1H, d, J 15.6 Hz), 5.81 (1H, d, J 8.8 Hz), and 3.53 (3H, s).

Anal. Calcd. for C₂₅H₁₉Cl₂N₃O₂:

C, 64.67; H, 4.12; N, 9.05.

Found: C, 64.28; H, 4.24; N, 8.83%.

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EXAMPLE 12

E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(4-methylphenyl)-2-propenamide

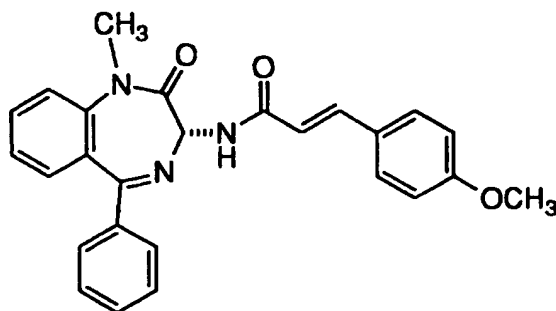
m.p. 133-135°C, $[\alpha]_D^{+90.4^\circ}$ (c=0.125, CH₂Cl₂).

δ_H (CDCl₃) 7.68-7.19 (15H, m), 6.59 (1H, d, J 15.6 Hz), 5.70 (1H, d, J 8.0 Hz), 3.50 (3H, s), and 2.38 (3H, s).

Anal. Calcd. for C₂₆H₂₃N₃O₂:

C, 76.26; H, 5.66; N, 10.26.

Found: C, 75.93; H, 5.82; N, 10.10%.

EXAMPLE 13

E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(4-methoxyphenyl)-2-propenamide

m.p. 129-133°C, $[\alpha]_D^{+89.9^\circ}$ (c 0.188, CH₂Cl₂).

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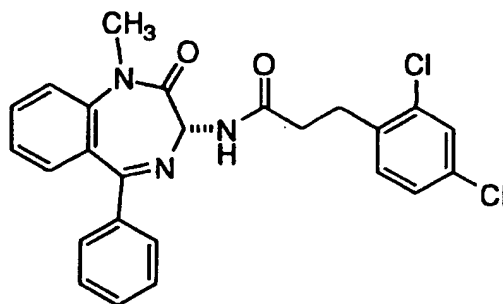
dH (CDCl₃) 7.65-7.24 (14H, m), 6.92 (1H, d, J 8.8 Hz), 6.50 (1H, d, J 15.6 Hz), 5.69 (1H, d, J 8.0 Hz), 3.84 (3H, s), and 3.50 (3H, s).

Anal. Calcd. for C₂₆H₂₃N₃O₃·0.30H₂O:

C, 72.48; H, 5.52; N, 9.75.

Found: C, 72.75; H, 5.60; N, 9.36%.

EXAMPLE 14



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

m.p. 92-95°C, [α]_D 90.5° (c = 0.196, CH₂Cl₂).

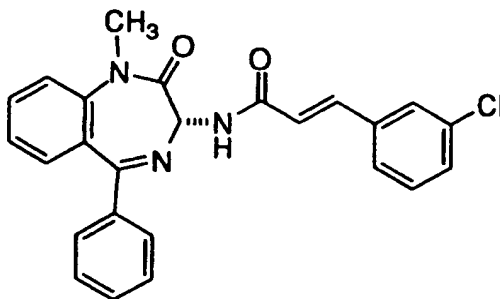
dH (CDCl₃) 7.62-7.15 (13H, m), 5.52 (1H, d, J 8.1 Hz), 3.47 (3H, s), 3.10 (2H, t, J 7.6 Hz), and 2.68 (2H, dd, J 7.6, 2.8 Hz).

Anal. Calcd. for C₂₅H₂₁Cl₂N₃O₂·0.20H₂O:

C, 63.89; H, 4.59; N, 8.94.

Found: C, 63.86; H, 4.62; N, 8.87%.

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EXAMPLE 15

E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(3-chlorophenyl)-2-propenamide

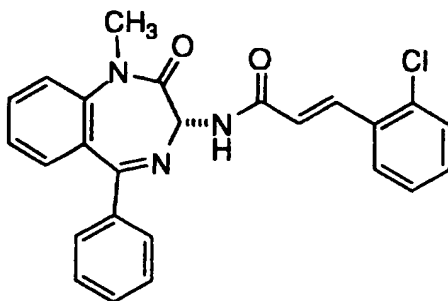
m.p. 229-231°C, $[\alpha]_D +86.2^\circ$ ($c = 0.225$, CH_2Cl_2).

^1H (CDCl_3) 7.64-7.26 (15H, m), 6.62 (1H, d, J 15.6 Hz), 5.66 (1H, d, J 8.1 Hz), and 3.50 (3H, s).

Anal. Calcd. for $\text{C}_{25}\text{H}_{20}\text{ClN}_3\text{O}_2$:

C, 69.85; H, 4.69; N, 9.77.

Found: C, 70.20; H, 4.83; N, 9.41%.

EXAMPLE 16

E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2-chlorophenyl)-2-propenamide

m.p. 128-131°C, $[\alpha]_D = +61.7^\circ$ ($c = 0.196$, CH_2Cl_2).

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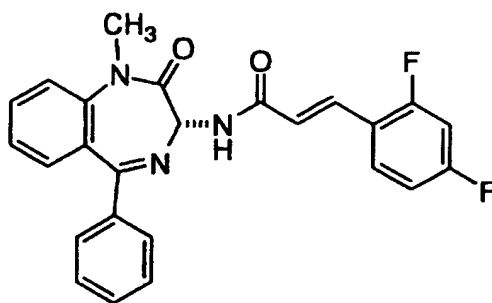
δ H (CDCl₃) 8.06 (1H, d, J 15.6 Hz), 7.65-7.28 (14H, m), 6.62, (1H, d, J 15.6 Hz), 5.68 (1H, d, J 8.3 Hz), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₂₀ClN₃O₂·0.20H₂O:

C, 69.27; H, 4.74; N, 9.69.

Found: C, 69.21; H, 4.68; N, 9.45%.

EXAMPLE 17



E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-difluorophenyl)-2-propenamide

m.p. 121-123°C, [α]_D +76.8° (c = 0.111, CH₂Cl₂).

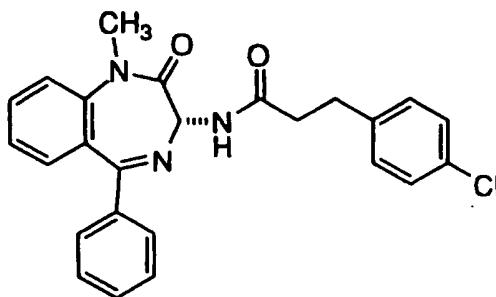
δ H (CDCl₃) 7.71 (1H, d, J 15.9 Hz), 7.64-7.24 (11H, m), 6.92-6.84 (2H, m), 6.69 (1H, d, J 15.9 Hz), 5.67 (1H, d, J 8.1 Hz), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₁₉F₂N₃O₂·0.10H₂O:

C, 69.31; H, 4.47; N, 9.70.

Found: C, 69.28; H, 4.57; N, 9.31%.

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EXAMPLE 18

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(4-chlorophenyl)propanamide

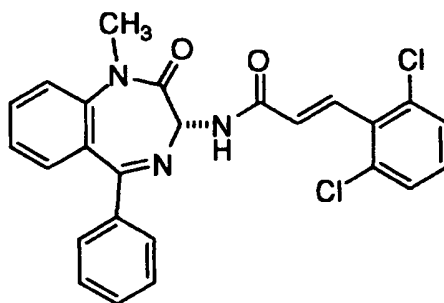
m.p. 203-205°C, $[\alpha]_D +99.2^\circ$ (c = 0.300, CH₂Cl₂).

¹H (CDCl₃) 7.62-7.16 (14H, m), 5.52 (1H, d, J 8.1 Hz), 3.47 (3H, s), 2.99 (2H, t, J 7.7 Hz), and 2.67 (2H, t, J 7.7 Hz).

Anal. Calcd. for C₂₅H₂₂ClN₃O₂:

C, 69.52; H, 5.13; N, 9.73.

Found: C, 69.50; H, 5.15; N, 9.72%.

EXAMPLE 19

E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,6-dichlorophenyl)-2-propenamide

m.p. 121-124°C, $[\alpha]_D +69.0^\circ$ (c = 0.342, CH₂Cl₂).

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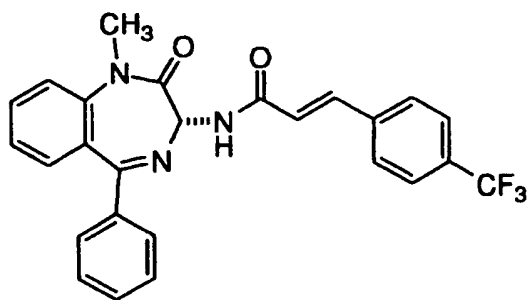
dH (CDCl₃) 7.79 (1H, d, J 16.1 Hz), 7.64-7.15 (13H, m), 6.78 (1H, d, J 15.8 Hz), 5.69 (1H, d, J 8.1 Hz), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₁₉Cl₂N₃O₂·0.15PhCH₃:

C, 65.44; H, 4.23; N, 8.79.

Found: C, 65.40; H, 4.38; N, 8.85%.

EXAMPLE 20



E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-[4-(trifluoromethyl)phenyl]-2-propenamide

m.p. 133-137°C, [α]_D +68.7° (c = 0.115, CH₂Cl₂).

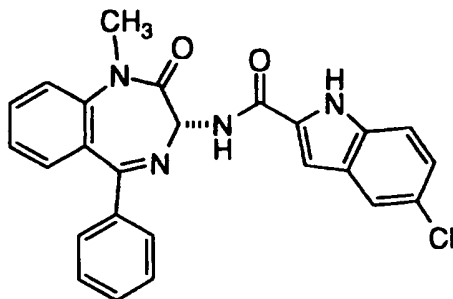
dH (CDCl₃) 7.72-7.25 (15H, m), 6.71 (1H, d, J 15.6 Hz), 5.67 (1H, d, J 8.1 Hz), and 3.51 (3H, s).

Anal. Calcd. for C₂₆H₂₀F₃N₃O₂:

C, 67.38; H, 4.35; N, 9.07.

Found: C, 67.38; H, 4.45; N, 8.95%.

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EXAMPLE 21

(+)-5-Chloro-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]indole-2-carboxamide

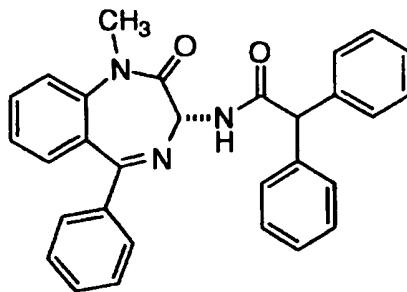
m.p. 160-164°C, $[\alpha]_D^{+103.8^\circ}$ (c = 0.160, CH₂Cl₂).

¹H (CDCl₃) 9.71 (1H, br s), 8.13 (1H, d, J 7.8 Hz), 7.68-7.09 (13H, m), 5.75 (1H, d, J 7.8 Hz), and 3.53 (3H, s).

Anal. Calcd. for C₂₅H₁₉ClN₄O₂·0.25H₂O·0.15PhCH₃:

C, 67.84; H, 4.49; N, 12.15.

Found: C, 67.80; H, 4.41; N, 12.07%.

EXAMPLE 22

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2,2-diphenylethanamide

m.p. 200-201°C, $[\alpha]_D^{+97.0^\circ}$ (c = 0.168, CH₂Cl₂).

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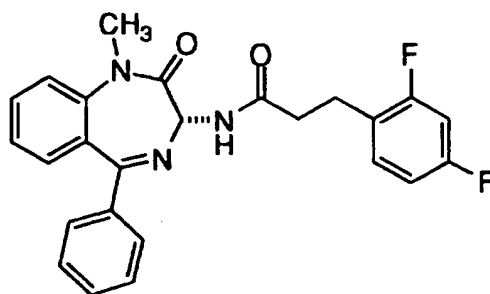
δ H (CDCl₃) 7.60-7.22 (20H, m), 5.58 (1H, d, J 8.1 Hz), 5.08 (1H, s), and 3.44 (3H, s).

Anal. Calcd. for C₃₀H₂₅N₃O₂·0.15PhCH₃:

C, 78.79; H, 5.55; N, 8.88.

Found: C, 78.81; H, 5.63; N, 9.07%.

EXAMPLE 23



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-difluorophenyl)propanamide

m.p. 79-81°C, [α]_D +92.9° (c = 0.105, CH₂Cl₂).

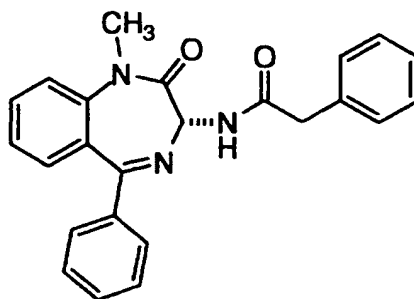
δ H (CDCl₃) 7.62-7.56 (3H, m), 7.50-7.19 (8H, m), 6.82-6.76 (2H, m), 5.52 (1H, d, J 8.1 Hz), 3.47 (3H, s), 3.01 (2H, t, J 7.6 Hz), and 2.69 (2H, m).

Anal. Calcd. for C₂₅H₂₁F₂N₃O₂:

C, 69.27; H, 4.88; N, 9.69.

Found: C, 68.96; H, 4.99; N, 9.47%.

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EXAMPLE 24

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-phenylethanamide

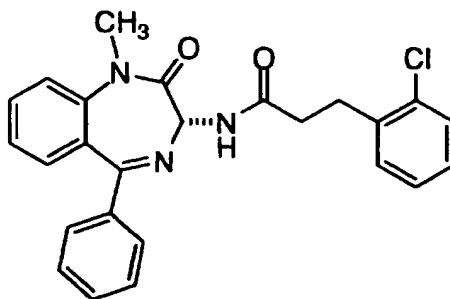
m.p. 241-242°C (dec.), $[\alpha]_D^{+85.5^\circ}$ (c = 0.159, CH₂Cl₂).

δ_H (CDCl₃) 7.59-7.55 (3H, m), 7.46-7.22 (12H, m), 5.51 (1H, d, J 8.1Hz), 3.72 (2H, s), and 3.44 (3H, s).

Anal. Calcd. for C₂₄H₂₁N₃O₂·0.55H₂O:

C, 73.28; H, 5.66; N, 10.68.

Found: C, 73.25; H, 5.38; N, 10.47%.

EXAMPLE 25

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2-chlorophenyl)propanamide

m.p. 158.5-159.5°C, $[\alpha]_D^{+95.8^\circ}$ (c = 0.224, CH₂Cl₂).

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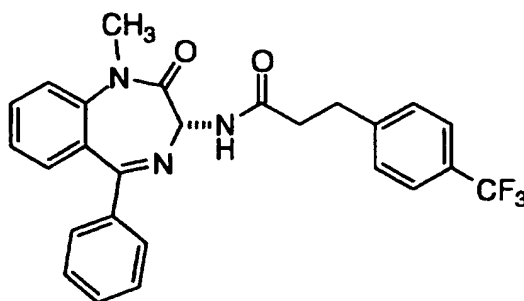
dH (CDCl₃) 7.62-7.57 (3H, m), 7.47-7.16 (11H, m), 5.55 (1H, d, J 8.1 Hz), 3.47 (3H, s), 3.14 (2H, t, J 7.9 Hz), and 2.75-2.69 (2H, m).

Anal. Calcd. for C₂₅H₂₂ClN₃O₂·0.15H₂O:

C, 69.09; H, 5.17; N, 9.67.

Found: C, 69.05; H, 5.12; N, 9.63%.

EXAMPLE 26



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-[4-(trifluoromethyl)phenyl]propanamide

m.p. 175-176°C, [α]_D +86.5° (c = 0.141, CH₂Cl₂).

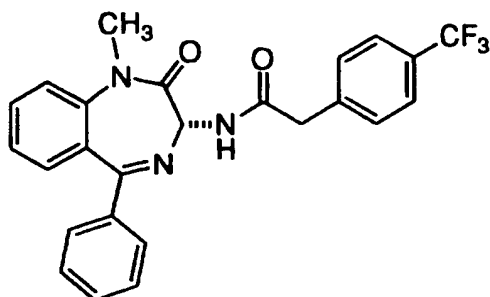
dH (CDCl₃) 7.62-7.54 (5H, m), 7.47-7.22 (9H, m), 5.52 (1H, d, J 8.1 Hz), 3.47 (3H, m), 3.08 (2H, t, J 7.6Hz), and 2.72 (2H, m).

Anal. Calcd. for C₂₆H₂₂F₃N₃O₂·0.80H₂O:

C, 65.08; H, 4.93; N, 8.76.

Found: C, 65.03; H, 4.63; N, 8.72%.

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EXAMPLE 27

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzo-
diazepin-3-yl]-2-[4-(trifluoromethyl)phenyl]ethanamide

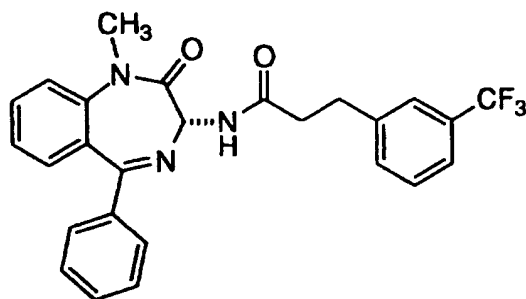
m.p. 224-226°C, $[\alpha]_D +68.0^\circ$ (c = 0.153, CH₂Cl₂).

¹H (CDCl₃) 7.63-7.55 (4H, m), 7.51-7.33 (8H, m), 7.26-7.23 (2H, m),
5.51 (1H, d, J 8.1 Hz), 3.77 (2H, s), and 3.46 (3H, s).

Anal. Calcd. for C₂₅H₂₀F₃N₃O₂:

C, 66.51; H, 4.47; N, 9.31.

Found: C, 66.46; H, 4.36; N, 9.10%.

EXAMPLE 28

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzo-
diazepin-3-yl]-3-[3-(trifluoromethyl)phenyl]propanamide

m.p. 135-136°C, $[\alpha]_D +78.8^\circ$ (c = 0.134, CH₂Cl₂).

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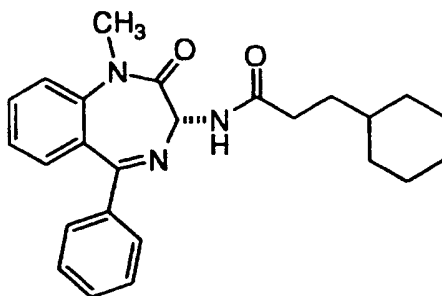
δ H (CDCl₃) 7.62-7.56 (3H, m), 7.49-7.22 (11H, m), 5.53 (1H, d, J 8.1 Hz), 3.47 (3H, s), 3.08 (2H, t, J 7.3 Hz), and 2.72 (2H, m).

Anal. Calcd. for C₂₆H₂₂F₃N₃O₂:

C, 67.09; H, 4.76; N, 9.03.

Found: C, 67.03; H, 4.73; N, 9.13%.

EXAMPLE 29



(+)-3-Cyclohexyl-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]propanamide

m.p. 144.5-145.5°C, [α]_D +83.1° (c = 0.116, CH₂Cl₂).

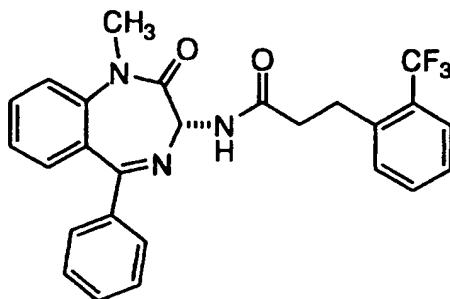
δ H (CDCl₃) 7.62-7.56 (3H, m), 7.46-7.21 (7H, m), 5.55 (1H, d, J 8.3 Hz), 3.48 (3H, s), 2.41-2.36 (2H, m), 1.77-1.58 (7H, m), 1.31-1.16 (4H, m), and 0.98-0.90 (2H, m).

Anal. Calcd. for C₂₅H₂₉N₃O₂:

C, 74.41; H, 7.24; N, 10.41.

Found: C, 74.46; H, 7.27; N, 10.58%.

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EXAMPLE 30

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-[2-(trifluoromethyl)phenyl]propanamide

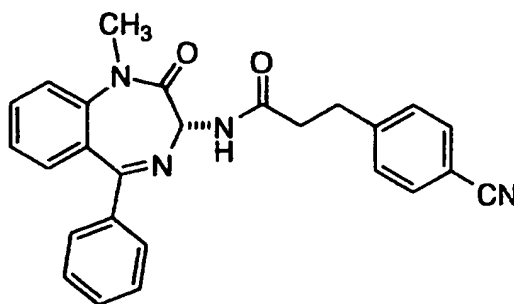
m.p. 110-113°C, $[\alpha]_D^{25} +79.2^\circ$ (c = 0.376, CH₂Cl₂).

¹H (CDCl₃) 7.65-7.57 (4H, m), 7.50-7.22 (10H, m), 5.55 (1H, d, J 8.0 Hz), 3.47 (3H, s), 3.20 (2H, t, J 7.9 Hz), and 2.70 (2H, dt, J 7.9, 3.3 Hz).

Anal. Calcd. for C₂₆H₂₂F₃N₃O₂:

C, 67.09; H, 4.76; N, 9.03.

Found: C, 66.97; H, 4.76; N, 8.93%.

EXAMPLE 31

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(4-cyanophenyl)propanamide

m.p. 81-85°C, $[\alpha]_D^{25} +91.0^\circ$ (c = 0.111, CH₂Cl₂).

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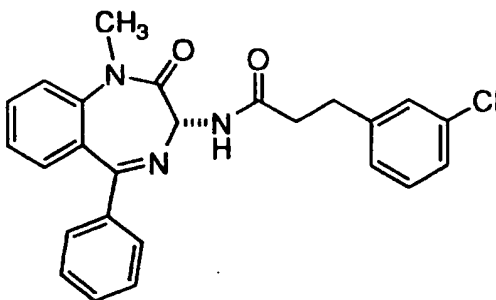
δ H (CDCl₃) 7.64-7.55 (4H, m), 7.48-7.16 (10H, m), 5.50 (1H, d, J 8.3 Hz), 3.47 (3H, s), 3.08 (2H, t, J 7.6 Hz), and 2.74-2.69 (2H, m).

Anal. Calcd. for C₂₆H₂₂N₄O₂·0.60H₂O·0.50PhCH₃:

C, 73.93; H, 5.62; N, 11.69.

Found: C, 73.98; H, 5.61; N, 11.71%.

EXAMPLE 32



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(3-chlorophenyl)propanamide

m.p. 157-159°C, [α]_D +90.7° (c = 0.134, CH₂Cl₂).

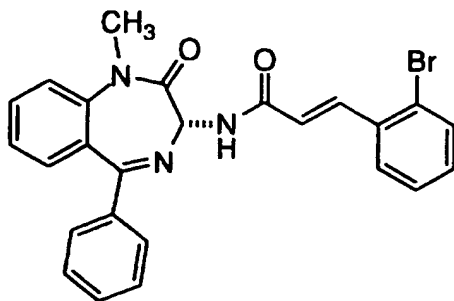
δ H (CDCl₃) 7.62-7.57 (3H, m), 7.47-7.12 (11H, m), 5.53 (1H, d, J 8.1 Hz), 3.47 (3H, s), 3.00 (2H, t, J 7.3 Hz), and 2.71-2.66 (2H, m).

Anal. Calcd. for C₂₅H₂₂ClN₃O₂·0.55H₂O:

C, 67.96; H, 5.27; N, 9.51.

Found: C, 67.99; H, 5.18; N, 9.26%.

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EXAMPLE 33

E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2-bromophenyl)-2-propenamide

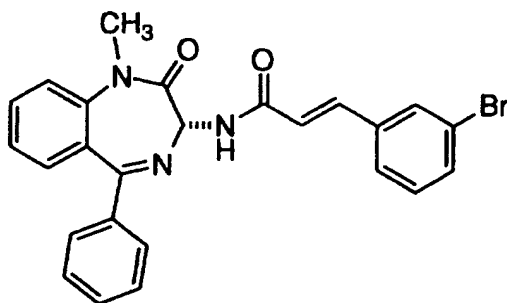
m.p. 113-116°C, $[\alpha]_D^{+44.2^\circ}$ (c = 0.113, CH₂Cl₂).

^1H (CDCl₃) 8.03 (1H, d, J 15.6 Hz), 7.64-7.16 (14H, m), 6.57 (1H, d, J 15.6 Hz), 5.68 (1H, d, J 8.1 Hz), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₂₀BrN₃O₂·0.60H₂O·0.30PhCH₃:

C, 63.48; H, 4.58; N, 8.19.

Found: C, 63.49; H, 4.38; N, 8.19%.

EXAMPLE 34

E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(3-bromophenyl)-2-propenamide

m.p. 221-223 d°C, $[\alpha]_D^{+65.5^\circ}$ (c = 0.206, CH₂Cl₂).

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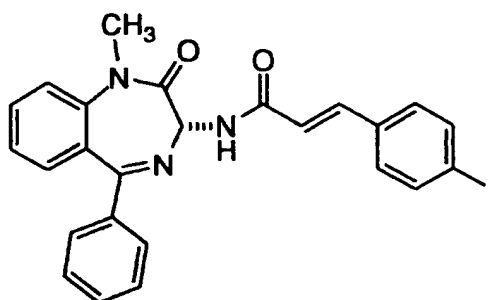
dH (CDCl₃) 7.69 (1H, br s), 7.64-7.57 (4H, m), 7.51-7.37 (6H, m), 7.29-7.19 (4H, m), 6.62 (1H, d, J 15.6 Hz), 5.66 (1H, d, J 8.1 Hz), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₂₀BrN₃O₂·0.35H₂O·0.20PhCH₃:

C, 63.54; H, 4.46; N, 8.42.

Found: C, 63.50; H, 4.39; N, 8.42%.

EXAMPLE 35



E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(4-iodophenyl)-2-propenamide

m.p. 137-140°C, [α]_D +67.9° (c = 0.268, CH₂Cl₂).

dH (CDCl₃) 7.75-7.72 (2H, m), 7.64-7.36 (8H, m), 7.29-7.16 (5H, m), 6.63 (1H, d, J 15.6 Hz), 5.66 (1H, d, J 8.1 Hz), and 3.50 (3H, m).

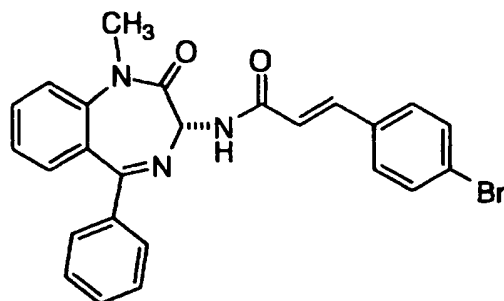
Anal. Calcd. for C₂₅H₂₀IN₃O₂·0.30PhCH₃:

C, 59.29; H, 4.06; N, 7.65.

Found: C, 59.29; H, 3.90; N, 7.40%.

EXAMPLE 36

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E-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(4-bromophenyl)-2-propenamide

m.p. 121-124°C, $[\alpha]_D^{25} +75.6^\circ$ (c = 0.201, CH₂Cl₂).

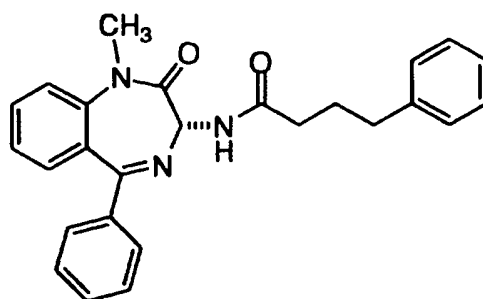
¹H (CDCl₃) 7.64-7.57 (3H, m), 7.55-7.35 (11H, m), 7.28-7.24 (1H, m), 6.62 (1H, d, J 15.6 Hz), 5.66 (1H, d, J 8.1 Hz), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₂₀BrN₃O₂:

C, 63.30; H, 4.25; N, 8.86.

Found: C, 63.50; H, 4.20; N, 8.78%.

EXAMPLE 37



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-4-phenylbutanamide

m.p. 65-74°C, $[\alpha]_D^{25} +77.4^\circ$ (c = 0.155, CH₂Cl₂).

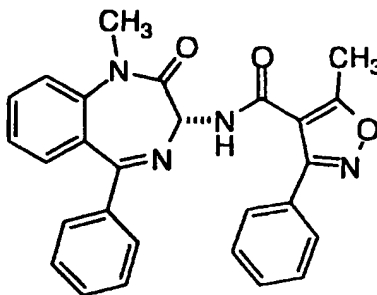
¹H (CDCl₃) 7.62-7.56 (3H, m), 7.46-7.19 (12H, m), 5.55 (1H, d, J 8.1 Hz), 3.47 (3H, s), 2.71 (2H, t, J 7.6 Hz), 2.42-2.37 (2H, m), and 2.09-2.01 (2H, m).

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Anal. Calcd. for $C_{26}H_{25}N_3O_2 \cdot 0.30H_2O$:

C, 74.91; H, 6.19; N, 10.08.

Found: C, 74.93; H, 6.05; N, 10.07%.

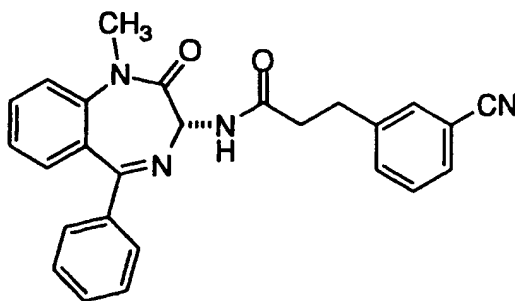
EXAMPLE 38

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-5-methyl-3-phenylisoxazole-4-carboxamide

m.p. 123-126°C, $[\alpha]_D^{+122.0^\circ}$ ($c = 0.199$, CH_2Cl_2). δ_H ($CDCl_3$) 7.79-7.76 (2H, m), 7.62-7.32 (11H, m), 7.26-7.21 (2H, m), 5.61 (1H, d, J 7.9 Hz), 3.42 (3H, s), and 2.76 (3H, s).Anal. Calcd. for $C_{27}H_{22}N_4O_3 \cdot 0.40H_2O$:

C, 70.85; H, 5.02; N, 12.24.

Found: C, 70.84; H, 4.91; N, 11.92%.

EXAMPLE 39

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(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(3-cyanophenyl)propanamide

m.p. 110-112°C, $[\alpha]_D^{+84.2^\circ}$ (c = 0.202, CH₂Cl₂).

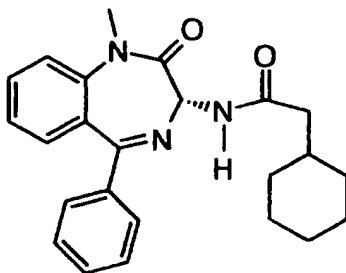
¹H (CDCl₃) 7.63-7.22 (14H, m), 5.51 (1H, d, J 8.1 Hz), 3.47 (3H, s), 3.06 (2H, t, J 7.8 Hz), and 2.74-2.68 (2H, m).

Anal. Calcd. for C₂₆H₂₂N₄O₂·0.50H₂O:

C, 72.37; H, 5.37; N, 12.98.

Found: C, 72.52; H, 5.12; N, 12.59%.

EXAMPLE 40



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]cyclohexanethanamide

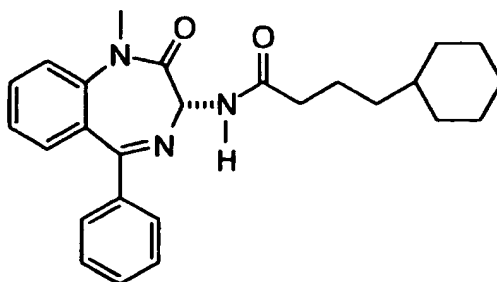
m.p. 144-146°C, $[\alpha]_D^{+72.1^\circ}$ (c=1.000, MeOH).

Anal. Calcd. for C₂₄H₂₇N₃O₂·0.20H₂O:

C, 73.33; H, 7.03; N, 10.69.

Found: C, 73.27; H, 7.02; N, 10.76%.

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EXAMPLE 41

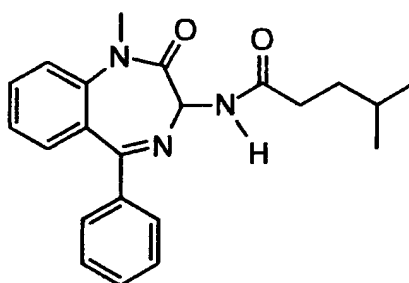
(+)-4-Cyclohexyl-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]butanamide

$[\alpha]_D +57.7^\circ$ ($c=0.440$, MeOH).

Anal. Calcd. for $C_{26}H_{31}N_3O_2$:

C, 74.79; H, 7.48; N, 10.06.

Found: C, 74.8; H, 7.78; N, 10.05%.

EXAMPLE 42

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-4-methylpentanamide

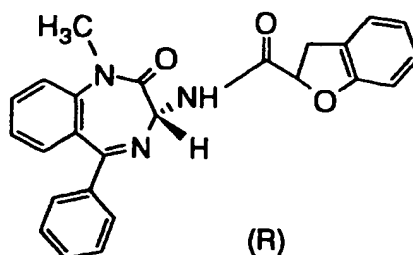
m.p. 123-125°C, $[\alpha]_D +66.8^\circ$ ($c=0.500$, MeOH).

Anal. Calcd. for $C_{22}H_{25}N_3O_2 \cdot 0.45H_2O$:

C, 71.12; H, 7.03; N, 11.31.

Found: C, 71.08; H, 6.81; N, 11.42%.

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EXAMPLE 43

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2,3-dihydrobenzofuran-2-carboxamide

Diisopropylethylamine (0.3 mL, 223 mg, 1.72 mmol) was added to a stirred, cooled (0 °C) solution of 3(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (*J. Org. Chem.* 1987, 52, 3232-3239) (400 mg, 1.5 mmol), 2,3-dihydrobenzofuran-2-carboxylic acid (274 mg, 1.7 mmol), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (583 mg, 3.0 mmol), and 1-hydroxybenzotriazole (479 mg, 3.1 mmol) in DMF (4.5 mL). The mixture was stirred at room temperature for 18 h., poured into aqueous hydrochloric acid (3M, 12 mL) and extracted with ethyl acetate (3 x 20 mL). The combined organic fractions were washed with saturated aqueous sodium hydrogen carbonate (20 mL) and brine (20 mL), dried (MgSO₄) and evaporated under reduced pressure. The residue was crystallized from 2-chloro-2-methylpropane/hexane to give (+)-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2,3-dihydrobenzofuran-2-carboxamide as a colorless solid (156 mg, 25%), m.p. 141-180°C, [α]_D +127.1° (c=0.425, CHCl₃).

dH (CDCl₃) (3:1 Mixture of diastereoisomers) 8.44 (1H, m), 7.65-6.91 (13H, m), 5.52 (1H, m), 5.28 (1H, m), and 3.70-3.40 (5H, m).

Anal. Calcd. for C₂₅H₂₁N₃O₃·0.25 Hexane

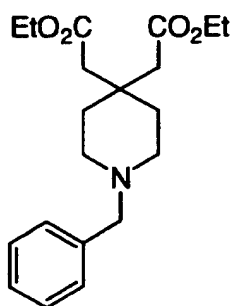
C, 73.50; H, 5.70; N, 9.71.

Found: C, 74.12; H, 5.57; N, 9.71%.

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EXAMPLE 44

(+)-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-1'-(1,1-dimethylethoxycarbonyl)spiro(cyclohexan-4,4'-piperidine)-1-carboxamide

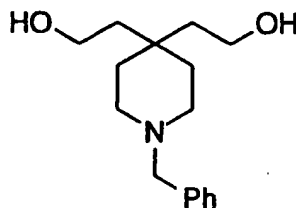
Step A:**Diethyl 1-benzylpiperidine-4,4-diacetate**

Ethanol (120 mL) was cooled in ice and ammonia bubbled through to give a saturated solution. 1-Benzyl-4-piperidone (40.0g, 211 mmol) and ethyl cyanoacetate (47.8g, 423 mmol) were added, the reaction vessel stoppered and stored at 0°C overnight. The solid was collected, washed with ethanol and ether and dried in vacuo to give a yellow solid (68.86g). The solid (58.86g) was dissolved in a mixture of sulfuric acid (70 mL, 98%) and water (60 mL) and heated under reflux for three days the mixture cooled and most of the water evaporated. The residue was azeotroped with ethanol (4x750 mL), further ethanol (500 mL) added and the mixture heated under reflux for 20h, cooled in ice and sodium carbonate (100g) added slowly with vigorous stirring. The ethanol was evaporated under reduced pressure, water (800 mL) added and the mixture extracted with methylene chloride (3x400 mL). The combined organic extracts were dried (Na₂SO₄) and the solvent evaporated to give diethyl 1-benzylpiperidine-4,4-diacetate (37.51g). A small portion of this was purified by flash column chromatography.

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NMR (300 MHz, CDCl_3) δ : 7.2-7.4 (m, 5H), 4.11 (q, $J=7.3\text{Hz}$, 4H), 3.50 (s, 2H), 2.56 (s, 4H), 2.4 (m, 4H), 1.7 (m, 4H), 1.24 (t, $J=7.3\text{Hz}$, 6H).

Step B:



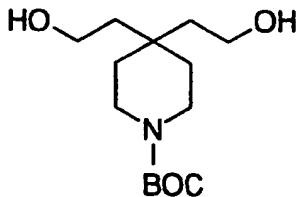
1-Benzylpiperidine-4,4-diethanol

A solution of the diester (12.2 g, 35 mmol) in ether (25 mL) was added to a cooled (-30°C) and stirred suspension of LiAlH_4 (2.1 g, 55 mmol) in ether (400 mL), under argon. THF (60 mL) was added and the reaction mixture allowed to warm to room temperature. After recooling to 0°C , water (2.2 mL), 1M NaOH (4.4 mL) and water (5 mL) were added, the reaction mixture stirred vigorously for 30 min and the solid filtered off, washing well with ether. The combined filtrates were evaporated to afford a white solid which was triturated with ether to give 8 g of 1-benzylpiperidine-4,4-diethanol.

m.p. $75-78^\circ\text{C}$

NMR (300 MHz, CDCl_3) δ : 7.2-7.4 (m, 5H), 3.7 (t, $J = 6.8\text{ Hz}$, 4H), 3.52 (s, 2H), 2.7 (brs, 2H), 2.43 (m, 4H), 1.66 (t, $J = 6.8\text{ Hz}$, 4H), 1.5 (m, 4H).

Step C:

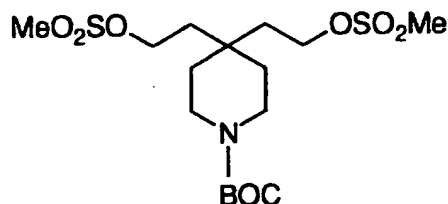


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1-t-Butoxycarbonylpiperidine-4,4-diethanol

The benzylamine (2.07 g, 7.9 mmol) was dissolved in methanol (60 mL), BOC₂O (1.72 g, 7.9 mmol) added and the mixture hydrogenated at 50 psi over 10% palladium hydroxide on charcoal (200 mg) for 18 hours. The reaction mixture was filtered through celite, washed with methanol and the filtrate evaporated to give 1-t-butoxycarbonylpiperidine-4,4-diethanol (2.0 g).

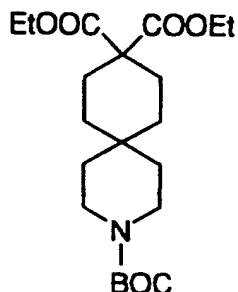
NMR (300 MHz, CDCl₃) d: 3.7 (m, 4H), d 3.3 (m, 6H), 1.65 (t, J = 6.8 Hz, 4H), 1.41 (s, 9H).

Step D:1-t-Butoxycarbonylpiperidine-4,4-diethanol, bis(methanesulfonate)

The diol (2.41 g, 8.9 mmol) was dissolved in dichloromethane (50 mL), the solution cooled to -20°C under argon before addition of triethylamine (3.7 mL, 26 mmol) and methanesulfonyl chloride (1.6 mL, 20 mmol). After 30 min., the reaction mixture was poured into ice cold 10% citric acid and extracted with ether (X3). The combined extracts were washed with water, saturated NaHCO₃ and brine, dried (MgSO₄) and the solvent evaporated to afford 1-t-butoxycarbonylpiperidine-4,4-diethanol, bis(methanesulfonate) (3.2g).

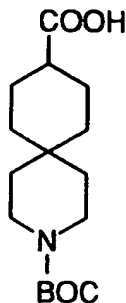
NMR (300 MHz, CDCl₃) d: 4.32 (t, J = 7.1 Hz, 4H), 3.4 (m, 4H), 3.04 (s, 6H), 1.89 (t, J = 7.1 Hz, 4H).

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Step E:Diethyl 3-t-butyloxycarbonyl-3-azaspiro[5.5]undecane-9,9-dicarboxylate

To a slurry of 60% NaH (2.04 g, 0.51 mole) in toluene (160 mL), under argon, was slowly added diethyl malonate (3.72 mL, 24.3 mmol). The mixture was cooled to 0°C and the bis-mesylate 1 (7.0 g, 16.3 mmol) added as a solid and the mixture heated to reflux for 18 hours. The reaction was quenched into 10% citric acid (100 mL) and the product extracted with CH₂Cl₂ (2x150 mL). The extracts were dried (Na₂SO₄), concentrated to an oil, and chromatographed on silica to give 3.83 g (60%) of diethyl 3-t-butyloxycarbonyl-3-azaspiro-[5.5]undecane-9,9-dicarboxylate.

¹H NMR (CDCl₃) δ: 1.22 (t, 6H), 1.4 (s, 9H), 2.0 (m, 4H), 3.35 (m, 4H), 4.2 (q, 4H).

Step F:

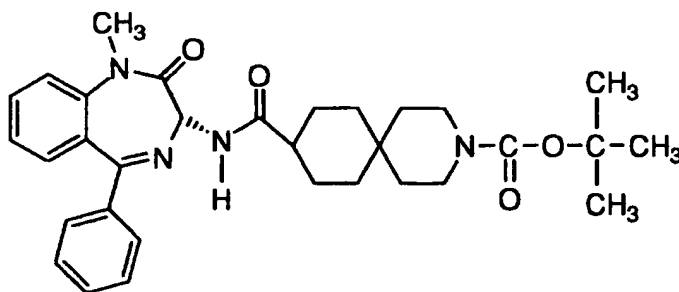
- 69 -

3-t-Butyloxycarbonyl-3-azaspiro[5.5]undecane-9-carboxylic acid

To a solution of the diester 2 (3.69 g, 0.0093 m) in THF (50 mL) was added 1N LiOH (47 mL). The reaction was stirred for 3 days at 25°C, diluted with water (50 mL) and pH adjusted to 2.2 with KHSO₄. The product was extracted into ethyl acetate (2x75 mL), dried (Na₂SO₄), and concentrated to a foam (3.5 g). The solid was melted in a flask at 140°C for 2 hours, cooled and the oil dissolved in THF (15 mL), 1N LiOH (10 mL) added and mixture stirred overnight at 30°C. The reaction was concentrated to remove THF, diluted with water (20 mL) and washed with diethyl ether (10 mL). The pH was adjusted to 2.5 with KHSO₄ and product extracted (3x50 mL) with ethyl acetate. The extracts were dried (Na₂SO₄), filtered and concentrated to yield 3-t-butyloxycarbonyl-3-azaspiro[5.5]undecane-9-carboxylic acid as a foam (2.48 g, 90%).

¹H NMR (CDCl₃, partial) δ: 1.45 (s, 9H), 3.4 (m, 4H).

Employing the procedure substantially as described in Example 43 but substituting an appropriate acid for the 2,3-dihydro-benzofuran-2-carboxylic acid, the following compounds were prepared:

Step G:

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-1'-(1,1-dimethylethoxycarbonyl)spiro(cyclohexan-4,4'-piperidine)-1-carboxamide

m.p. 135-138°C, [α]_D +58.8° (C=0.925, CHCl₃).

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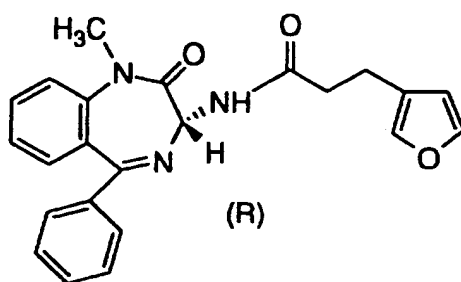
dH (CDCl₃) 7.61-7.23 (10H, m), 5.54 (1H, d, J 9.0 Hz), 3.47 (3H, s), 3.37 (4H, m), 2.28 (1H, m), and 1.81-1.18 (21H, s).

Anal. Calcd. for C₃₂H₄₀N₄O₄:

C, 70.56; H, 7.40; N, 10.29.

Found: C, 70.21; H, 7.40; N, 10.16%.

EXAMPLE 45



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(furan-2-yl)propanamide

m.p. 115-118°C, [α]_D +65.8° (c=0.800, CHCl₃).

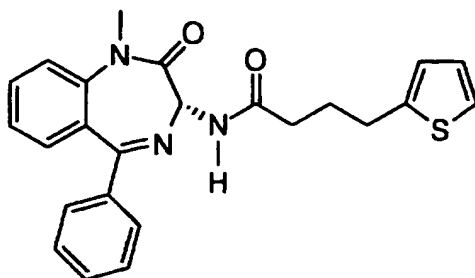
dH (CDCl₃) 7.62-7.26 (11H, m), 6.28 (1H, dd, J 3.2, 2.0 Hz), 6.08 (1H, dd, J 3.2, 0.7 Hz), 5.58 (1H, d, J 8.1 Hz), 3.48 (3H, s), 3.04 (2H, t, J 7.6 Hz), and 2.75 (2H, m).

Anal. Calcd. for C₂₃H₂₁N₃O₃·0.3Hexane:

C, 72.07; H, 6.15; N, 10.17.

Found: C, 71.78; H, 6.30; N, 9.77%.

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EXAMPLE 46

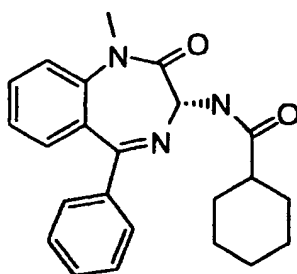
(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzo-
diazepin-3-yl]-4-(2-thienyl)butanamide

m.p. 170-180°C, $[\alpha]_D +63.5^\circ$ (c=1.000, MeOH).

Anal. Calcd. for $C_{24}H_{23}N_3O_2S \cdot 0.95H_2O$:

C, 66.32; H, 5.77; N, 9.67.

Found: C, 66.32; H, 5.34; N, 9.40%.

EXAMPLE 47

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzo-
diazepin-3-yl]cyclohexylcarboxamide

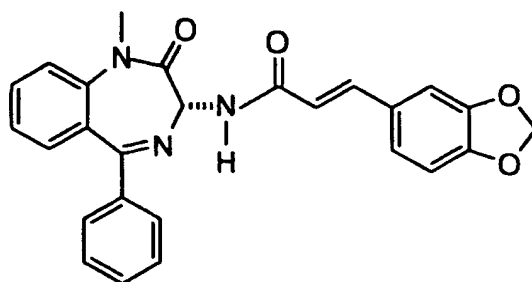
m.p. 213-214°C, $[\alpha]_D +62.4^\circ$ (c=1.000, MeOH).

Anal. Calcd. for $C_{23}H_{24}N_3O_2$:

C, 73.77; H, 6.46; N, 11.22.

Found: C, 73.86; H, 6.81; N, 11.15%.

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EXAMPLE 48

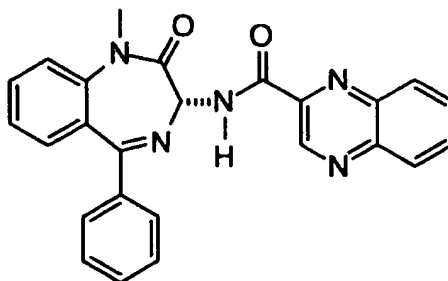
(E)-(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(3,4-methylenedioxyphenyl)-2-propenamide

m.p. 143-145°C, $[\alpha]_D +62.3^\circ$ (c=0.960, MeOH).

Anal. Calcd. for $C_{25}H_{21}N_3O_4 \cdot 0.10H_2O \cdot 0.20Et_2O$:

C, 69.78; H, 5.27; N, 9.46.

Found: C, 69.78; H, 4.98; N, 9.28%.

EXAMPLE 49

(+)-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-quinoxalinecarboxamide

$[\alpha]_D +85.8^\circ$ (c=0.360, MeOH).

Anal. Calcd. for $C_{25}H_{19}N_5O_2$:

C, 69.96; H, 4.90; N, 15.33.

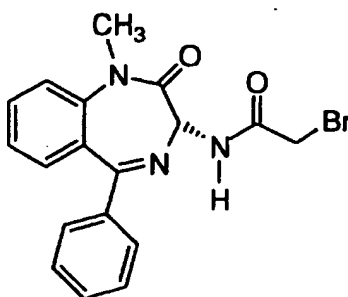
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Found: C, 69.95; H, 4.72; N, 15.25%.

EXAMPLE 50

(+)-N-[(3R)-2,3-Dihydro-2-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-(phenylamino)acetamide

Step A:

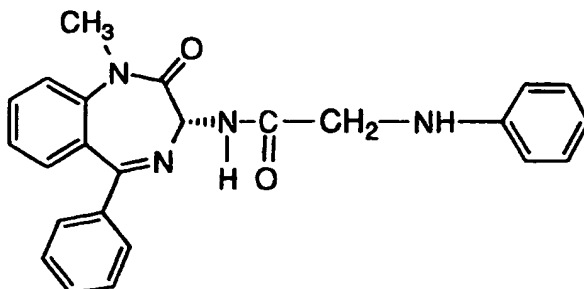


N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-bromoacetamide

Bromoacetyl bromide (165 mL, 383 mg, 1.9 mmol) was added to an ice cooled solution of 3(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (*J. Org. Chem.* 1987, 52, 3232-3239) (500 mg, 1.88 mmol) and triethylamine (264 mL, 192 mg, 1.9 mmol) in methylene chloride (10 mL) and the mixture was stirred at room temperature for 1 h. The mixture was washed with water (3 x 10 mL), dried (MgSO₄) and the solvent was evaporated under reduced pressure to give N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-bromoacetamide as a colorless foam (760 mg, 100%).

¹H (CDCl₃) 8.24 (1H, d, J 7.8 Hz), 7.64-7.24 (9H, m), 5.48 (1H, d, J 7.8 Hz), 4.00 (2H, m), and 3.50 (3H, s).

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Step B:

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-(phenylamino)acetamide

Aniline (297 mL, 304 mg, 3.26 mmol) was added to a solution of N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-bromoacetamide (600 mg, 1.55 mmol) in ethanol (25 mL) and the mixture was heated under reflux for 24 h. The mixture was cooled and the solid was collected and recrystallized from ethanol (20 mL) to give (+)-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-(phenylamino)acetamide as a colorless solid (500 mg, 81%), m.p. 245-246°C, $[\alpha]_D^{+119}$ (C=0.850, CHCl₃).

δ_H (CDCl₃) 8.26 (1H, d, J 8.3 Hz), 7.63-7.20 (12H, m), 6.81 (1H, t, J 7.3 Hz), 6.72 (2H, d, J 7.6 Hz), 5.56 (1H, d, J 8.3 Hz), 3.95 (2H, d, J 1.5 Hz), and 3.45 (3H, s).

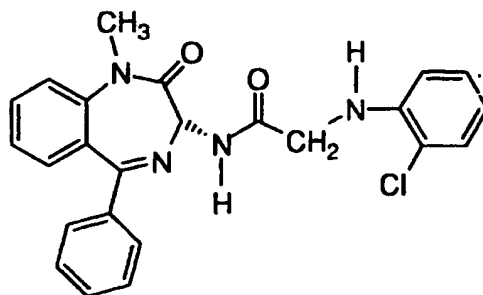
Anal. Calcd. for C₂₄H₂₂N₄O₂:

C, 72.34; H, 5.57; N, 14.06.

Found: C, 72.37; H, 5.59; N, 14.32%.

Employing the procedure substantially as described above, but substituting 2-chloroaniline or 4-(trifluoromethyl)aniline for the aniline, the following compounds were prepared:

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EXAMPLE 51

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-(2-chlorophenylamino)acetamide

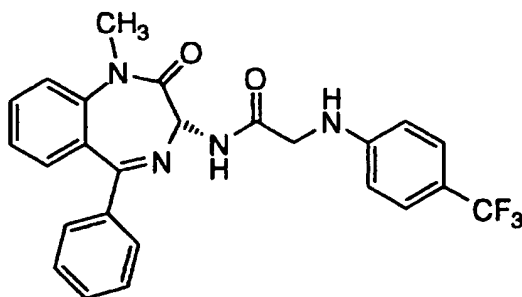
m.p. 222-224°C, $[\alpha]_D^{+111}$ (c=0.973, CHCl₃).

^1H (CDCl₃) 8.15 (1H, d, J 8.3 Hz), 7.60-7.16 (12H, m), 6.71 (2H, m), 5.57 (1H, d, J 8.3 Hz), 4.01 (2H, d, J 2.7 Hz), and 3.45 (3H, s).

Anal. Calcd. for C₂₄H₂₁ClN₄O₂:

C, 66.59; H, 4.89; N, 12.94.

Found: C, 66.40; H, 4.94; N, 12.92%.

EXAMPLE 52

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-[4-(trifluoromethyl)phenylamino]acetamide

m.p. 218-219°C, $[\alpha]_D^{+91.9}$ (c = 0.419, CHCl₃).

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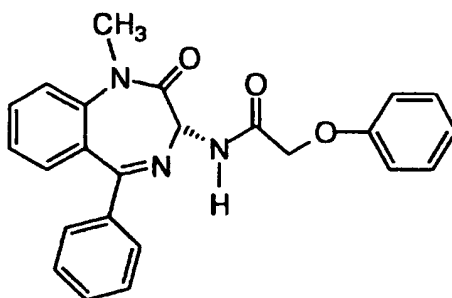
dH (CDCl₃) 8.13 (1H, d, J 9.0 Hz), 7.70-7.25 (12H, m), 6.72 (2H, d, J 8.7 Hz), 5.60 (1H, d, J 9.0 Hz), 4.05 (2H, m), and 3.50 (3H, s).

Anal. Calcd. for C₂₅H₂₁F₃N₄O₂·0.7H₂O:

C, 62.68; H, 4.71; N, 11.69.

Found: C, 62.47; H, 4.32; N, 11.44%.

EXAMPLE 53



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-(phenoxy)acetamide

Phenol (104 mg, 1.1 mmol) was added to a suspension of sodium hydride (60% dispersion in mineral oil, 44 mg, 1.1 mmol) in toluene (10 mL). When hydrogen evolution had stopped, N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-bromoacetamide (400 mg, 1.04 mmol) was added and the mixture was stirred at room temperature for 18 h. The mixture was washed with water (3 x 15 mL), dried (MgSO₄) and the solvent was evaporated under reduced pressure. The residue was triturated with 2-propanol and the solid was collected and recrystallized from 2-propanol (5 mL) to give (+)-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-(phenoxy)acetamide as a colorless solid (112 mg, 27%), m.p. 126-128°C, [α]_D +81.6 (C=0.692, CHCl₃).

dH (CDCl₃) 8.49 (1H, d, J 8.2 Hz), 7.64-7.01 (14H, m), 5.61 (1H, d, J 8.2 Hz), 4.65 (1H, d, J 14.6 Hz), 4.58 (1H, d, J 14.6 Hz), and 3.50 (3H, s).

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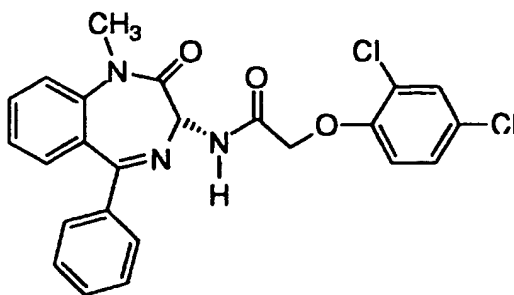
Anal. Calcd. for C₂₄H₂₁N₃O₃:

C, 72.17; H, 5.30; N, 10.52.

Found: C, 71.84; H, 5.25; N, 10.41%.

Employing the procedure substantially as described above, but substituting 2,4-dichlorophenol, thiophenol or 2,4-dichlorothiophenol for the phenol, the following compounds were prepared:

EXAMPLE 54



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-(2,4-dichlorophenoxy)acetamide

m.p. 206°C, [α]_D +31.1° (c=0.289, CHCl₃).

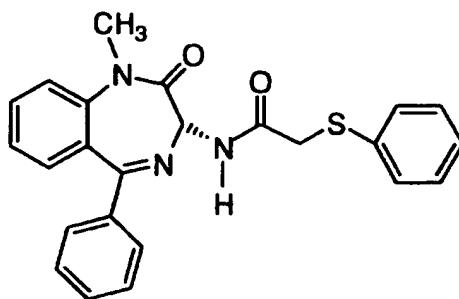
¹H (CDCl₃) 8.75 (1H, d, J 9.0 Hz), 7.65-7.20 (11H, m), 6.90 (1H, d, J 8.7 Hz), 5.60 (1H, d, J 9.0 Hz), 4.65 (2H, m), and 3.50 (3H, s).

Anal. Calcd. for C₂₄H₁₉Cl₂N₃O₃·0.3H₂O:

C, 60.85; H, 4.17; N, 8.87.

Found: C, 60.80; H, 4.04; N, 8.87%.

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EXAMPLE 55

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-(phenylthio)acetamide

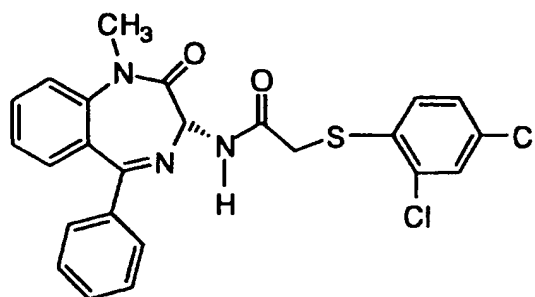
$[\alpha]_D +104.9^\circ$ (c=0.316, CHCl₃).

δ_H (CDCl₃) 8.50 (1H, d, J 9.0 Hz), 7.60-7.20 (14H, m), 5.50 (1H, d, J 9.0 Hz), 3.75 (2H, m), and 3.45 (3H, s).

Anal. Calcd. for C₂₄H₂₁N₃O₂S:

C, 69.37; H, 5.10; N, 10.11.

Found: C, 68.98; H, 5.06; N, 9.76%.

EXAMPLE 56

(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-2-(2,4-dichlorophenylthio)acetamide

$[\alpha]_D +97.4^\circ$ (c=0.286, CHCl₃).

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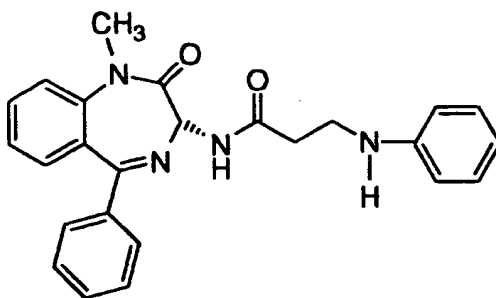
δ H (CDCl₃) 8.35 (1H, d, J 9.0 Hz), 7.70-7.20 (12H, m), 5.50 (1H, d, J 9.0 Hz), 3.70 (2H, m), and 3.50 (3H, s).

Anal. Calcd. for C₂₄H₁₉Cl₂N₃O₂S:

C, 59.51; H, 3.95; N, 8.67.

Found: C, 59.32; H, 3.95; N, 8.65%.

EXAMPLE 57



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(phenylamino)propanamide

3-Bromopropionyl chloride (2.01 mL, 3.428 g, 20 mmol) was added to an ice cooled solution of 3(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (*J. Org. Chem.* 1987, 52, 3232-3239) (5.0 g, 18.8 mmol) and triethylamine (2.79 mL, 2.02 g, 20 mmol) in methylene chloride (85 mL) and the mixture was stirred at room temperature for 18 h. The mixture was washed with saturated aqueous sodium hydrogen carbonate (85 mL), water (2 x 85 mL), and brine (85 mL), dried (MgSO₄) and the solvent was evaporated under reduced pressure. A sample (0.5 g, 1.25 mmol) was dissolved in ethanol (25 mL), aniline (230 mL, 233 mg, 2.5 mmol) was added and the mixture was heated under reflux for 70 h. The mixture was cooled and the solid was collected and recrystallized from ethanol to give (+)-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(phenylamino)propanamide as a colorless solid, m.p. 218-221°C, [α]_D +58.2° (c=0.585, CHCl₃).

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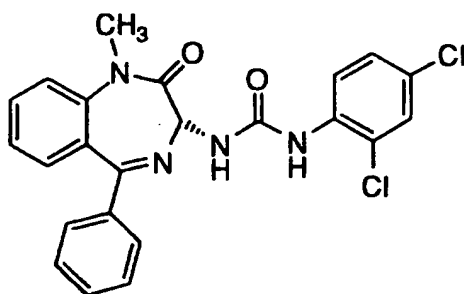
δ H (CDCl₃) 7.60-6.71 (16H, m), 5.54 (1H, d, J 8.1 Hz), 3.54 (2H, t, J 6.1 Hz), 3.52 (3H, s), and 2.70 (2H, m).

Anal. Calcd. for C₂₅H₂₄N₄O₂·0.5EtOH:

C, 71.70; H, 6.25; N, 12.87.

Found: C, 71.42; H, 5.98; N, 12.84%.

EXAMPLE 58



(+)-1-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)urea

2,4-Dichlorophenyliisocyanate (188 mg, 1.0 mmol) was added to a solution of 3(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (*J. Org. Chem.* 1987, 52, 3232-3239) (265 mg, 1.0 mmol) in tetrahydrofuran (20 mL). The mixture was stirred at room temperature for 18 h. and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with CH₂Cl₂/MeOH (99.5:0.5) and the residue was crystallized from CH₂Cl₂/hexane to give (+)-1-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)urea as a colorless solid, m.p. 215-216.5°C, [α]_D +76.2° (c=0.261, CHCl₃).

δ H (CDCl₃) 8.10 (1H, d, J 9.0 Hz), 7.65-6.95 (13H, m), 5.50 (1H, d, J 9.0 Hz), and 3.50 (3H, s).

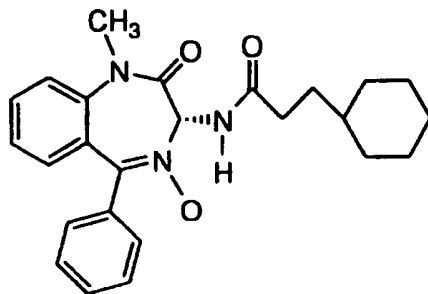
Anal. Calcd. for C₂₃H₁₈Cl₂N₄O₂·0.3H₂O:

C, 60.22; H, 4.09; N, 12.21.

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Found: C, 60.28; H, 3.89; N, 12.10%.

EXAMPLE 59



(-)-3-Cyclohexyl-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-4-oxido-5-phenyl-1H-1,4-benzodiazepin-3-yl]propanamide

3-Chloroperoxybenzoic acid (80%, 0.32 g, 1.5 mmol) was added to a solution of (+)-3-cyclohexyl-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]propanamide (0.60 g, 1.5 mmol) in dichloromethane (25 mL) and the mixture was stirred at room temperature for 18 h. Further 3-chloroperoxybenzoic acid (80%, 0.1 g, 0.5 mmol) was added and the mixture was stirred for 24 h. The mixture was washed with saturated aqueous sodium hydrogen carbonate (4 x 25 mL), water (2 x 25 mL) and brine (25 mL), dried (MgSO₄) and the solvent was evaporated under reduced pressure. The residue was recrystallized from toluene/hexane (65:35) to give (-)-3-cyclohexyl-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-4-oxido-5-phenyl-1H-1,4-benzodiazepin-3-yl]propanamide as colorless prisms, m.p. 222-224°C, [α]_D -80.7° (c=1.15, CHCl₃).

¹H (CDCl₃) 7.71-7.23 (10H, m), 6.01 (1H, d, J 9.3 Hz), 3.54 (3H, s), 2.48 (2H, m), and 1.76-0.89 (13H, m).

Anal. Calcd. for C₂₅H₂₉N₃O₃·0.5H₂O:

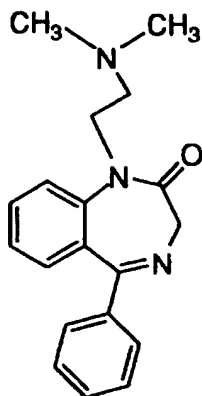
C, 70.06; H, 7.06; N, 9.81.

Found: C, 70.10; H, 6.80; N, 9.79%.

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EXAMPLE 60

N-[2,3-Dihydro-1-(2-dimethylaminoethyl)-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

Step A:

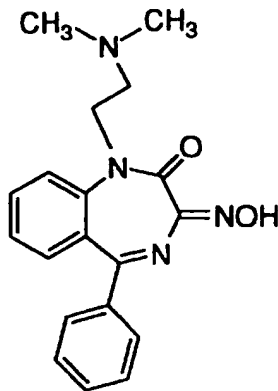
2,3-Dihydro-1-(2-dimethylaminoethyl)-5-phenyl-1H-1,4-benzodiazepin-2-one

2,3-Dihydro-5-phenyl-1H-1,4-benzodiazepin-2-one (1.00 g, 4.23 mmol) was added to hexane washed sodium hydride (60% dispersion in mineral oil, 186 mg, 4.65 mmol) in DMF (5 mL). Further DMF (10 mL) was added and the mixture was stirred at room temperature. 2-(Dimethylamino)ethyl chloride hydrochloride (0.73 g, 5 mmol) was added to hexane washed sodium hydride (60% dispersion in mineral oil, 200 mg, 5.0 mmol) in DMF (5 mL) and the mixtures were combined. Potassium iodide (1 crystal) was added and the mixture was stirred at 110°C for 30 min. The solvent was evaporated under reduced pressure, water was added and the mixture was extracted with ethyl acetate. The combined organic fractions were washed with water (2 x), dried (MgSO₄) and the solvent was evaporated under reduced pressure to give 2,3-dihydro-1-(2-dimethylaminoethyl)-5-phenyl-1H-1,4-benzodiazepin-2-one (1.21 g, 93%).

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δ H (CDCl₃) 7.63-7.16 (9H, m), 4.77 (1H, d, J 10.6 Hz), 4.41 (1H, m), 3.80 (1H, m), 3.78 (1H, d, J 10.6 Hz), 2.49 (2H, m), and 2.13 (6H, s).

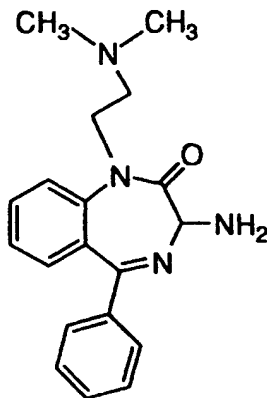
Step B:



2,3-Dihydro-1-(2-dimethylaminoethyl)-3-hydroxyimino-5-phenyl-1H-1,4-benzodiazepin-2-one

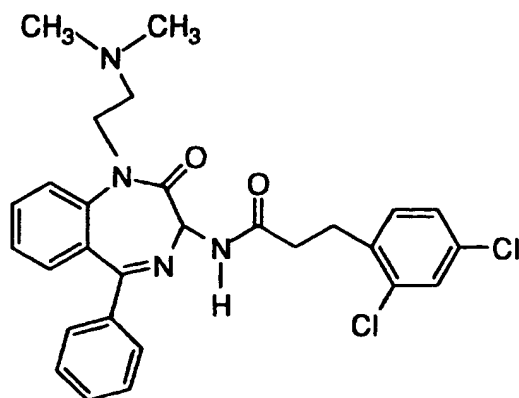
2,3-Dihydro-1-(2-dimethylaminoethyl)-5-phenyl-1H-1,4-benzodiazepin-2-one (1.21 g, 3.9 mmol) was dissolved in toluene (20 mL). The mixture was cooled to -78 °C and potassium *t*-butoxide (1.0M solution in *t*-butanol, 4.72 mL, 4.72 mmol) was added. The mixture was stirred at -78 °C for 20 min., then isoamyl nitrite (0.63 mL, 0.55 g, 4.72 mmol) was added. The mixture was stirred at -78°C for 90 min. then allowed to warm to room temperature and poured into aqueous citric acid (1M, 10 mL). The pH was adjusted to 5.0 with aqueous sodium hydroxide then to 7.0 with saturated aqueous sodium hydrogen carbonate. The mixture was extracted with ethyl acetate (50 mL) and the organic layer was aged at room temperature. The solid which formed was collected and dried *in vacuo* to give 2,3-dihydro-1-(2-dimethylaminoethyl)-3-hydroxyimino-5-phenyl-1H-1,4-benzodiazepin-2-one (0.876 g, 66%) as a solid, m.p. 232-234°C. δ H (d₆-DMSO) 10.90 (1H, s), 7.72-7.25 (9H, m), 4.40 (1H, m), 3.80 (1H, m), 2.50 (2H, m), and 1.85 (6H, s).

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Step C:**3-Amino-2,3-dihydro-1-(2-dimethylaminoethyl)-5-phenyl-1H-1,4-benzodiazepin-2-one**

Ethyl isocyanate (320 mL, 287 mg, 4.0 mmol) was added to a mixture of 2,3-dihydro-1-(2-dimethylaminoethyl)-3-hydroxyimino-5-phenyl-1H-1,4-benzodiazepin-2-one (0.91 g, 2.7 mmol) and triethylamine (0.56 mL, 0.41 g, 4.0 mmol) in THF (30 mL). The mixture was heated under reflux for 7 h., further ethyl isocyanate (167 mL, 150 mg, 2.1 mmol) was added and the mixture was heated under reflux for 12 h. The mixture was cooled, the solvent was evaporated under reduced pressure and ethyl acetate (75 mL) and water (25 mL) were added. The organic phase was washed with water (4 x 25 mL), dried (MgSO₄) and evaporated under reduced pressure. The residue was dissolved in ethanol (100 mL), palladium on carbon (10%, 100 mg) was added and the mixture was shaken under hydrogen (50 p.s.i.) for 4.5 h. Further palladium on carbon (10%, 100 mg) was added and the mixture was shaken under hydrogen (50 p.s.i.) for 1.5 h. The mixture was filtered and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with CH₂Cl₂/MeOH to give 3-amino-2,3-dihydro-1-(2-dimethylaminoethyl)-5-phenyl-1H-1,4-benzodiazepin-2-one (180 mg, 17%).
dH (CDCl₃) 7.75-7.17 (9H, m), 4.45 (1H, s), 4.40 (1H, m), 3.82 (1H, m), 2.47 (4H, m), and 2.08 (6H, s).

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Step E:**N-[2,3-Dihydro-1-(2-dimethylaminoethyl)-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide**

Triethylamine was added to a mixture of 3-amino-2,3-dihydro-1-(2-dimethylaminoethyl)-5-phenyl-1H-1,4-benzodiazepin-2-one (180 mg, 0.6 mmol), 3-(2,4-dichlorophenyl)propanoic acid (131 mg, 0.6 mmol), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (115 mg, 0.6 mmol) and 1-hydroxybenzotriazole (81 mg, 0.6 mmol) in DMF (15 mL) until the pH was 9.0. The mixture was stirred at room temperature for 72 h. The solvent was evaporated under reduced pressure and ethyl acetate was added. The mixture was washed with water, saturated aqueous sodium hydrogen carbonate and water, dried (MgSO₄) and evaporated under reduced pressure. The residue was triturated with acetone and recrystallized from i-PrOH/MeOH to give N-[2,3-dihydro-1-(2-dimethylaminoethyl)-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)-propanamide as a solid, m.p. 199-201°C.

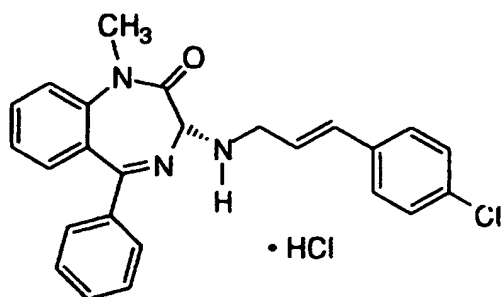
¹H (CDCl₃) 7.60-7.15 (13H, m), 5.50 (1H, d, J 8.0 Hz), 4.40 (1H, m), 3.80 (1H, m), 3.10 (2H, t, J 7.5 Hz), 2.70 (2H, t, J 7.5 Hz), 2.40 (2H, m), and 2.05 (6H, s).

Anal. Calcd. for C₂₈H₂₈Cl₂N₄O₂:

C, 64.25; H, 5.39; N, 10.70.

Found: C, 64.23; H, 5.40; N, 10.61%.

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EXAMPLE 61

(+)-3(R)-{N-[3-(4-chlorophenyl)prop-1-en-3-yl]amino}-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one hydrochloride

A mixture of 3(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (*J. Org. Chem.* 1987, 52, 3232-3239) (265 mg, 1 mmol), *E*-1-chloro-4-(3-chloro-1-propenyl)benzene (281 mg, 1.5 mmol), potassium carbonate (276 mg, 2 mmol) and potassium iodide (25 mg, 0.15 mmol) in acetonitrile (2 mL) was heated under reflux for 4 h. The mixture was cooled and poured into ethyl acetate (10 mL) and water (5 mL). The layers were separated and the aqueous layer was extracted with ethyl acetate (5 mL). The combined organic fractions were washed with brine, dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with EtOAc/Hexane (65:35 increasing to 100:0). The first compound to elute was suspended in ethanol (1 mL) and ethanolic HCl (6 M, 0.11 mL) was added. The mixture was stirred, then the solvent was evaporated under reduced pressure. The residue was triturated with ether and the solid was collected and dried *in vacuo* to give (+)-3(R)-{N,N-bis[1-(4-chlorophenyl)propen-3-yl]amino}-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one hydrochloride (235 mg, 39%) as a tan solid, m.p. 138-145°C, [α]_D +9.2° (c=0.500, MeOH).

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dH (d₆-DMSO) 11.2 (1H, br s), 7.77-7.31 (17H, m), 6.85 (2H, br m), 6.54 (2H, m), 5.20 (1H, br s), 4.60-4.00 (4H, m), and 3.46 (3H, s).

Anal. Calcd. for C₃₄H₂₉Cl₂N₃O.HCl.0.10EtOH:

C, 67.60; H, 5.08; N, 6.92.

Found: C, 67.60; H, 5.03; N, 7.03%.

The second compound to elute was suspended in ethanol (0.5 mL) and ethanolic HCl (6 M, 0.035 mL) was added. The mixture was stirred, then the solvent was evaporated under reduced pressure. The residue was triturated with ether and the solid was collected and dried in vacuo to give (+)-3(R)-{N-[3-(4-chlorophenyl)propen-3-yl]amino}-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one hydro-chloride (56 mg, 12%) as a yellow solid, m.p. 156-162°C, [α]_D +35° (c=0.100, MeOH).

dH (d₆-DMSO) 10.3 (1H, br s), 10.0 (1H, br s), 7.79-7.34 (13H, m), 6.78 (1H, d, J 15.9 Hz), 6.40 (1H, dt, J_d 15.9, J_t 9.0 Hz), 5.13 (1H, s), 4.00 (2H, m), and 3.46 (3H, s).

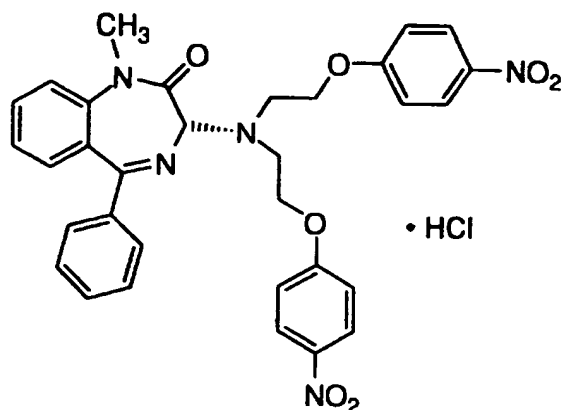
Anal. Calcd. for C₂₅H₂₂ClN₃O.HCl.0.10EtOH.0.40H₂O:

C, 65.20; H, 5.30; N, 9.05.

Found: C, 65.14; H, 5.09; N, 9.33%.

Employing the procedure substantially as described above, but substituting 1-(2-bromoethoxy)-4-nitrobenzene or 4-chlorobenzene-propanol methanesulfonate for the E-1-chloro-4-(3-chloro-1-propenyl)-benzene, the following compounds were prepared:

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EXAMPLE 62

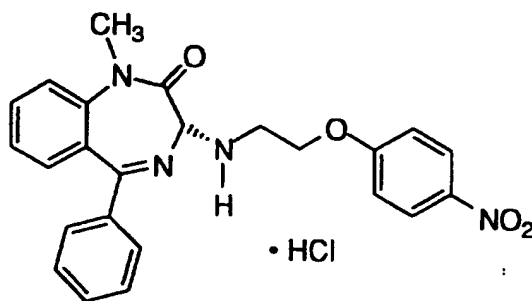
(+)-3(S)-{N,N-Bis[2-(4-nitrophenoxy)ethyl]amino}-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one hydrochloride
m.p. 126-145°C. $[\alpha]_D^{25} +5.0^\circ(0.100, \text{CHCl}_3)$.

^1H (d₆-DMSO) 8.20 (4H, d, J 9.2 Hz), 7.75-7.36 (9H, m), 7.08 (4H, d, J 9.2 Hz), 4.90 (1H, br s), 4.50 (4H, br s), 4.30-3.60 (5H, br m), and 3.34 (3H, s).

Anal. Calcd. for $\text{C}_{32}\text{H}_{29}\text{N}_5\text{O}_7 \cdot \text{HCl} \cdot 0.15\text{EtOH}$:

C, 60.71; H, 4.87; N, 10.96.

Found: C, 60.70; H, 4.87; N, 10.70%.

EXAMPLE 63

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(+)-3(R)-{N-[3-(4-Nitrophenoxy)ethyl]amino}-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one hydrochloride

m.p. 154-160°C, $[\alpha]_D^{25} +84.6^\circ$ (0.500, MeOH).

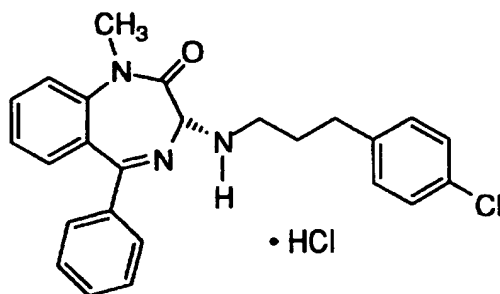
^1H NMR (d_6 -DMSO) 10.2 (1H, br s), 8.25 (2H, d, J 9.0 Hz), 7.83-7.41 (9H, m), 7.09 (2H, d, J 9.0 Hz), 5.21 (1H, s), 4.57 (2H, m), 3.70 (2H, m), 3.47 (3H, s), and 3.40 (1H, m).

Anal. Calcd. for $\text{C}_{24}\text{H}_{22}\text{N}_4\text{O}_4 \cdot \text{HCl} \cdot 0.15\text{EtOH} \cdot 0.20\text{H}_2\text{O}$:

C, 61.13; H, 5.13; N, 11.74.

Found: C, 61.12; H, 4.92; N, 11.64%.

EXAMPLE 64



(+)-3(R)-{N-[3-(4-Chlorophenyl)prop-1-yl]amino}-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one hydrochloride

m.p. 167-168°C, $[\alpha]_D^{25} +20.8^\circ$ (c=0.500, MeOH).

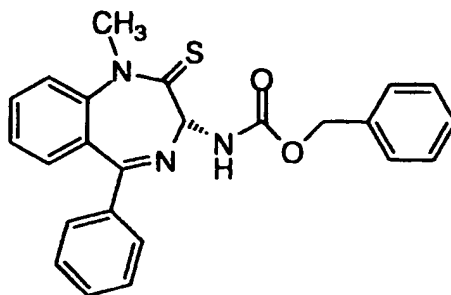
^1H NMR (d_6 -DMSO) 9.9 (2H, br m), 7.78-7.26 (13H, m), 5.08 (1H, s), 3.45 (3H, s), 3.20 (1H, m), 3.00 (1H, m), 2.70 (2H, t, J 7.4 Hz), and 2.05 (2H, m).

Anal. Calcd. for $\text{C}_{25}\text{H}_{24}\text{ClN}_3\text{O} \cdot \text{HCl}$:

C, 66.08; H, 5.55; N, 9.25.

Found: C, 65.81; H, 5.49; N, 9.30%.

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EXAMPLE 65

(+)-Phenylmethyl N-[(3R)-2,3-dihydro-1-methyl-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl]carbamate

A mixture of (+)-phenylmethyl N-[(3R)-2,3-dihydro-1-methyl-5-phenyl-2-oxo-1H-1,4-benzodiazepin-3-yl]carbamate (4.0 g, 10 mmol) and 2,4-bis(4-methoxyphenyl)-1,3-dithia-2,4-diphosphetane-2,4-disulfide (4.5 g, 11 mmol) in toluene (100 mL) was heated under reflux for 75 min. The mixture was cooled and the volume was reduced to 30 mL by evaporation under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with EtOAc/Hexane (75:25) to give (+)-phenylmethyl N-[(3R)-2,3-dihydro-1-methyl-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl]carbamate as a solid, m.p. 128-131°C, $[\alpha]_D^{25} +22.5^\circ$ ($c=0.656$, CHCl_3).
 ^1H (CDCl₃) 7.65-7.26 (15H, m), 5.50 (1H, d, J 8.8 Hz), 5.14 (2H, s), and 3.86 (3H, s).

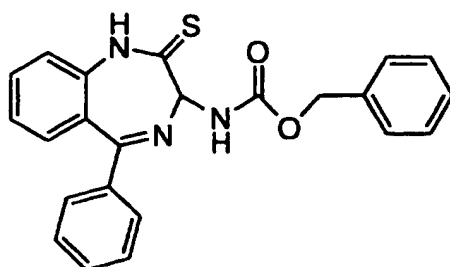
Anal. Calcd. for C₂₄H₂₁N₃O₂S.0.25H₂O:

C, 68.63; H, 5.16; N, 10.01.

Found: C, 68.28; H, 5.21; N, 10.06%.

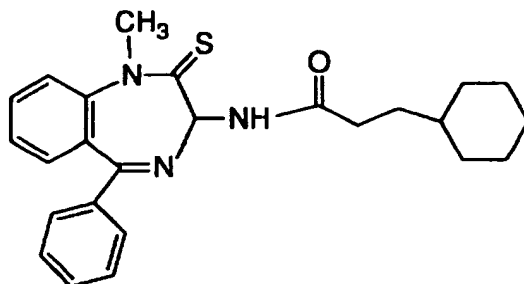
Employing the procedure substantially as described above, but substituting phenylmethyl N-[2,3-dihydro-5-phenyl-2-oxo-1H-1,4-benzodiazepin-3-yl]carbamate for the (+)-phenylmethyl N-[(3R)-2,3-dihydro-1-methyl-5-phenyl-2-oxo-1H-1,4-benzodiazepin-3-yl]carbamate, the following compound was prepared:

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EXAMPLE 66

Phenylmethyl N-[2,3-dihydro-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl]carbamate

dH (d₆-DMSO) 10.85 (1H, s), 8.42 (1H, d, J 8.6 Hz), 7.65-7.10 (14H, m), 5.10 (2H, s), and 5.05 (1H, d, J 8.6 Hz).

EXAMPLE 67

3-Cyclohexyl-N-(2,3-dihydro-1-methyl-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl)propanamide

Hydrogen bromide was bubbled at room temperature through a solution of (+)-phenylmethyl N-[(3R)-2,3-dihydro-1-methyl-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl]carbamate (0.9 g, 2.1 mmol), acetic acid (5 mL) and dichloromethane (5 mL). After 2 h., the solvent was evaporated under reduced pressure, ether was added and the

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solid was collected and dried in vacuo. A sample (0.58 g, 1.8 mmol) was suspended in THF (10 mL), triethylamine (0.24 mL, 0.18 g, 1.8 mmol) was added and the mixture was stirred at room temperature for 3 h. In a separate flask, oxalyl chloride (0.20 mL, 0.29 g, 2.3 mmol) was added to a solution of cyclohexanepropionic acid (0.33 mL, 0.30 g, 1.9 mmol) and DMF (1 drop) in THF (10 mL) and the mixture was stirred at room temperature for 3 h. The two mixtures were combined, triethylamine (0.32 mL, 0.23 g, 2.3 mmol) was added and the mixture was stirred at room temperature for 2.5 h. The solvent was evaporated under reduced pressure, water was added and the mixture was extracted with ethyl acetate. The combined organic fractions were washed with water, saturated aqueous sodium hydrogen carbonate, water (2 x) and brine, dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with CH₂Cl₂/MeOH (99.5:0.5) and the residue was recrystallized from EtOAc/Hexane to give 3-cyclohexyl-N-(2,3-dihydro-1-methyl-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl)propanamide as a solid, m.p. 219-221°C.

¹H (CDCl₃) 7.95 (1H, br d, J 8.6 Hz), 7.65-7.30 (9H, m), 5.72 (1H, d, J 8.6 Hz), 3.87 (3H, s), 2.41 (2H, t, J 7.6 Hz), and 1.80-0.85 (13H, m).

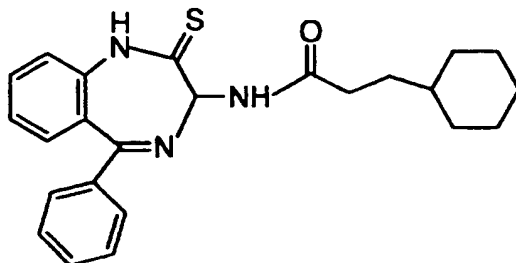
Anal. Calcd. for C₂₅H₂₉N₃OS.0.25H₂O:

C, 70.81; H, 7.01; N, 9.91.

Found: C, 70.80; H, 6.91; N, 9.95%.

Employing the procedure substantially as described above, but substituting phenylmethyl N-[2,3-dihydro-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl]carbamate for the (+)-phenylmethyl N-[(3R)-2,3-dihydro-1-methyl-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl]carbamate and an appropriate acid for the cyclohexanepropionic acid, the following compounds were prepared:

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EXAMPLE 68

3-Cyclohexyl-N-(2,3-dihydro-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl)propanamide

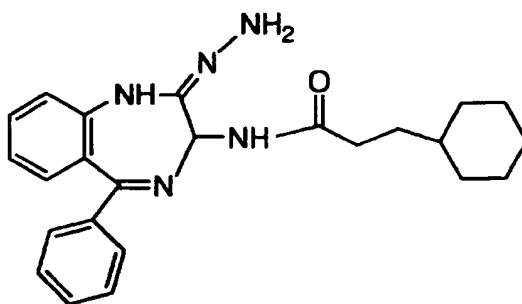
m.p. 113-119°C.

¹H (CDCl₃) 9.8 (1H, br s), 7.75-7.25 (10H, m), 5.75 (1H, d, J 8.1 Hz), 2.41 (2H, m), and 1.80-0.85 (13H, m).

Anal. Calcd. for C₂₄H₂₇N₃OS.0.8CH₂Cl₂:

C, 62.91; H, 6.09; N, 8.87.

Found: C, 62.88; H, 5.70; N, 9.12%.

EXAMPLE 69

3-Cyclohexyl-N-(2,3-dihydro-2-hydrazono-5-phenyl-1H-1,4-benzodiazepin-3-yl)propanamide

Hydrazine (53 mL, 56 mg, 1.8 mmol) was added to a solution of 3-cyclohexyl-N-(2,3-dihydro-1-methyl-5-phenyl-2-thioxo-

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1H-1,4-benzodiazepin-3-yl)propanamide (120 mg, 0.25 mmol) in methanol (3 mL). The mixture was stirred at room temperature for 3 h. and the solvent was evaporated under reduced pressure. Ethyl acetate was added and the mixture was washed with water and brine, dried (Na_2SO_4) and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (99.5:0.5 increasing to 98:2) to give 3-cyclohexyl-N-(2,3-dihydro-2-hydrazono-5-phenyl-1H-1,4-benzodiazepin-3-yl)propanamide as a foam.

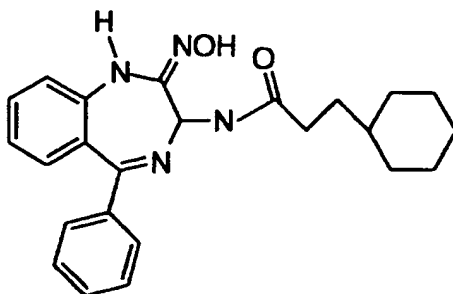
^1H (CDCl_3) 7.55-7.00 (11H, m), 5.75 (1H, d, J 7.6 Hz), 3.50 (2H, br s), 2.37 (2H, t, J 8.0 Hz), and 1.80-0.85 (13H, m).

Anal. Calcd. for $\text{C}_{24}\text{H}_{29}\text{N}_5\text{O} \cdot 0.8\text{CH}_3\text{OH} \cdot 0.15\text{CH}_2\text{Cl}_2$:

C, 67.82; H, 7.41; N, 15.85.

Found: C, 67.79; H, 7.46; N, 16.05%.

EXAMPLE 70



(E)- and (Z)-3-Cyclohexyl-N-(2,3-dihydro-2-hydroxyimino-5-phenyl-1H-1,4-benzodiazepin-3-yl)propanamide

A mixture of 3-cyclohexyl-N-(2,3-dihydro-1-methyl-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl)propanamide (740 mg, 1.83 mmol), hydroxylamine hydrochloride (140 mg, 2 mmol) and triethylamine (280 mL, 203 mg, 2 mmol) in methanol (15 mL)/THF (15 mL) was stirred at room temperature for 3 h. The solvent was evaporated under reduced pressure and the residue was purified by flash column

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chromatography on silica gel, eluting with CH₂Cl₂/MeOH (98:2). The residue recrystallized from ethyl acetate. The first isomer to crystallize was recrystallized from ethyl acetate to give (E)-3-cyclohexyl-N-(2,3-dihydro-2-hydroxyimino-5-phenyl-1H-1,4-benzodiazepin-3-yl)propanamide as a solid, m.p. 196°C.

¹H (d₆-DMSO) 12.20 (1H, s), 9.00 (1H, d, J 8.0 Hz), 7.70-7.30 (10H, m), 5.45 (1H, d, J 8.0 Hz), 2.30 (2H, m), and 1.80-0.75 (13H, m).

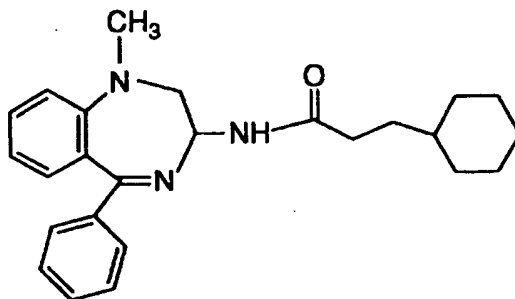
The second isomer to crystallize was recrystallized from methanol to give (Z)-3-cyclohexyl-N-(2,3-dihydro-2-hydroxyimino-5-phenyl-1H-1,4-benzodiazepin-3-yl)propanamide as a solid, m.p. 219°C. ¹H (d₆-DMSO) 9.95 (1H, s), 8.95 (1H, s), 8.75 (1H, d, J 8.0 Hz), 7.50-7.00 (9H, m), 5.70 (1H, d, J 8.0 Hz), 2.25 (2H, m), and 1.75-0.75 (13H, m).

Anal. Calcd. for C₂₄H₂₈N₄O₂:

C, 71.26; H, 6.98; N, 13.85.

Found: C, 70.89; H, 6.99; N, 13.55%.

EXAMPLE 71



3-Cyclohexyl-N-(2,3-dihydro-1-methyl-5-phenyl-1H-1,4-benzodiazepin-3-yl) propanamide

Freshly prepared Raney nickel (400 mg) was added to a solution of 3-cyclohexyl-N-(2,3-dihydro-1-methyl-5-phenyl-2-thioxo-1H-1,4-benzodiazepin-3-yl)propanamide (200 mg, 0.5 mmol) in ethanol (20 mL) and the mixture was stirred at room temperature for 2 h. The

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mixture was filtered and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with CH₂Cl₂/MeOH (99.75:0.25) to give 3-cyclohexyl-N-(2,3-dihydro-1-methyl-5-phenyl-1H-1,4-benzodiazepin-3-yl)propanamide as a foam.

¹H (CDCl₃) 7.60-6.80 (9H, m), 6.37 (1H, br d, J 6.6 Hz), 5.53 (1H, m), 3.60 (2H, m), 2.77 (3H, s), 2.21 (2H, t, J 8.0 Hz), and 1.85-0.80 (13H, m).

Anal. Calcd. for C₂₅H₃₁N₃O.0.2CH₂Cl₂:

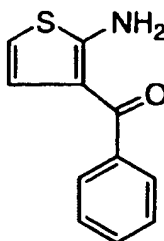
C, 74.45; H, 7.79; N, 10.34.

Found: C, 74.68; H, 7.87; N, 10.23%.

EXAMPLE 72

1-(2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-thieno-[2,3-e]-1,4-diazepin-3-yl)-3-(3-methyl-phenyl)urea

Step A:



(2-Amino-3-thienyl)phenylmethanone

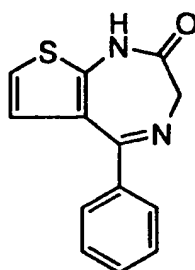
Triethylamine (6.8 mL, 4.94 g, 49 mmol) was added to a heated (33°C) mixture of b-oxobenzene propanenitrile (18.6 g, 128 mmol) and 1,2-dithiane-2,5-diol (9.8 g, 64 mmol) in ethanol (120 mL) and the mixture was stirred at 50°C for 18 h. The mixture was cooled and the solvent was evaporated under reduced pressure.

Dichloromethane was added, the mixture was washed with aqueous hydrochloric acid (0.5M), aqueous sodium hydroxide (1M) and brine,

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dried (Na_2SO_4) and the solvent was evaporated under reduced pressure. The residue was recrystallized from acetonitrile (150 mL) to give (2-amino-3-thienyl)-phenylmethanone as an orange solid (5.7 g, 44%). δ_{H} (CDCl_3) 7.70-7.35 (5H, m), 6.95 (2H, br s), 6.90 (1H, d, J 6.3 Hz), and 6.15 (1H, d, J 6.3 Hz).

Step B:



2,3-Dihydro-5-phenyl-1H-thien[2,3-e]-1,4-diazepin-2-one

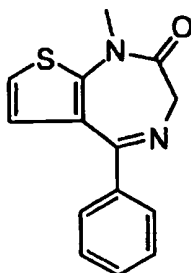
A solution of 1,3-dihydro-1,3-dioxo-2H-isoindole-2-acetyl chloride (8.6 g, 38 mmol) in dichloromethane (20 mL) was added slowly to a cooled (0°C) mixture of (2-amino-3-thienyl)phenylmethanone (6.8 g, 33 mmol), pyridine (6.34 mL, 6.20 g, 78 mmol) and 4-dimethylamino-pyridine (0.79 g, 6.5 mmol) in dichloromethane (130 mL). The mixture was stirred at 0°C for 30 min., diluted with dichloromethane (80 mL) and washed with aqueous hydrochloric acid (1M), saturated aqueous sodium hydrogen carbonate and brine. The mixture was dried (Na_2SO_4) and the solvent was evaporated under reduced pressure. The residue was triturated with ethanol and the solid was collected and dried *in vacuo* to give N-(3-benzoylthien-2-yl)-1,3-dihydro-1,3-dioxo-2H-isoindole-2-acetamide as a solid (9.8 g, 76%).

A mixture of N-(3-benzoylthien-2-yl)-1,3-dihydro-1,3-dioxo-2H-isoindole-2-acetamide (10.9 g, 28 mmol) and hydrazine (1.9 mL, 1.94 g, 60 mmol) in THF (500 mL) was heated under reflux for 4 h. The mixture was cooled, filtered and the solvent was evaporated under reduced pressure. Saturated aqueous sodium hydrogen carbonate was added and the mixture was extracted with ethyl acetate. The

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combined organic fractions were washed with brine, dried (Na_2SO_4) and the solvent was evaporated under reduced pressure. Acetic acid (300 mL) was added and the mixture was heated under reflux for 15 min. The mixture was cooled and the solvent was evaporated under reduced pressure. Saturated aqueous sodium hydrogen carbonate was added and the mixture was extracted with ethyl acetate. The combined organic fractions were washed with brine, dried (Na_2SO_4) and the solvent was evaporated under reduced pressure to give 2,3-dihydro-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one as a foam (3.5 g, 52%). δ_{H} (CDCl_3) 9.75 (1H, br s), 7.90-7.30 (5H, m), 6.87 (1H, d, J 6.0 Hz), 6.82 (1H, d, J 6.0 Hz), and 4.45 (2H, s).

Step C:



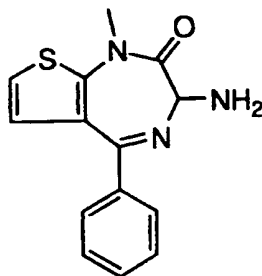
2,3-Dihydro-1-methyl-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one

Sodium hydride (60% dispersion in mineral oil, 757 mg, 11.3 mmol) was added to a cooled (0°C) solution of 2,3-dihydro-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one (2.61 g, 10.8 mmol) in DMF (7 mL). Further DMF (10 mL) was added and the mixture was stirred for 30 min. A solution of iodomethane (0.67 mL, 1.53 g, 10.8 mmol) in ether (20 mL) was added and the mixture was stirred for 1 h. The mixture was poured into water and the mixture was extracted with ethyl acetate. The combined organic fractions were washed with brine, dried (Na_2SO_4) and evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with

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CH₂Cl₂/MeOH (95:5) to give 2,3-dihydro-1-methyl-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one (1.5 g, 54%).
dH (CDCl₃) 7.67-7.35 (5H, m), 7.00 (1H, d, J 6.0 Hz), 6.85 (1H, d, J 6.0 Hz), 4.45 (2H, br s), and 3.50 (3H, s).

Step D:



3-Amino-2,3-dihydro-1-methyl-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one

2,3-Dihydro-1-methyl-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one (1.5 g, 5.8 mmol) was dissolved in toluene (30 mL). The mixture was cooled to -10°C and potassium t-butoxide (1.7 g, 15.1 mmol) was added. The mixture was stirred at -10°C for 15 min., then isoamyl nitrite (1.0 mL, 0.87 g, 7.4 mmol) was added. The mixture was stirred at -10°C for 1 h. then allowed to warm to room temperature and poured into water (50 mL) and acetic acid (3 mL). The mixture was extracted with ethyl acetate and the combined organic fractions were washed with brine, dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with EtOAc/Hexane to give 2,3-dihydro-1-methyl-3-hydroxyimino-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one (0.80 g, 48%).

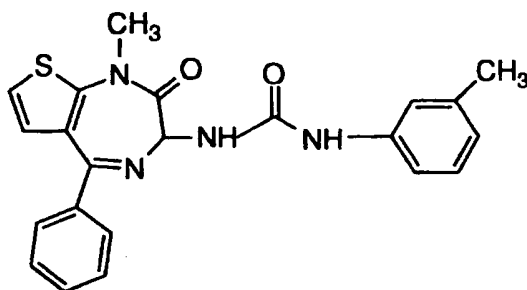
2,3-Dihydro-1-methyl-3-hydroxyimino-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one (0.80 g, 2.8 mmol) was dissolved in ethanol (40 mL) and Raney nickel (2 g) was added. The mixture was shaken under hydrogen (50 p.s.i.) for 5 days, adding further Raney nickel (10 g) in portions. The mixture was filtered and the solvent was evaporated

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under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with CH₂Cl₂/MeOH to give 3-amino-2,3-dihydro-1-methyl-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one (248 mg, 33%).

¹H (CDCl₃) 7.50-7.30 (5H, m), 7.05 (1H, d, J 6.0 Hz), 6.85 (1H, d, J 6.0 Hz), 4.57 (1H, s), 3.55 (3H, s), and 1.70 (2H, br s).

Step E:



1-(2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-3-yl)-3-(3-methylphenyl)urea

3-Methylphenylisocyanate (60 mL, 62 mg, 0.46 mmol) was added to a solution of 3-amino-2,3-dihydro-1-methyl-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one (124 mg, 0.46 mmol) in tetrahydrofuran (5 mL). The mixture was stirred at room temperature for 2 h. and the solvent was evaporated under reduced pressure. The residue was crystallized from EtOAc (4 mL) to give 1-(2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-3-yl)-3-(3-methylphenyl)urea as a solid (94 mg, 50%).

m.p. 128-130°C.

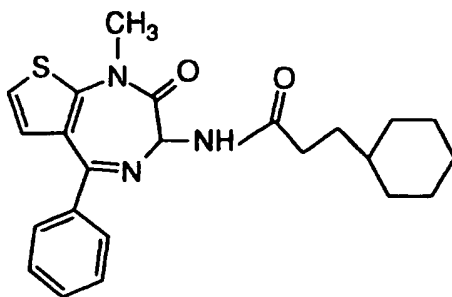
¹H (CDCl₃) 8.70 (1H, s), 7.65-6.75 (12H, m), 5.55 (1H, d, J 9.0 Hz), 3.55 (3H, s), and 2.30 (3H, s).

Anal. Calcd. for C₂₂H₂₀N₄O₂S.0.25H₂O:

C, 64.62; H, 4.99; N, 13.70.

Found: C, 64.68; H, 4.96; N, 13.70%.

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EXAMPLE 733-Cyclohexyl-N-(2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-3-yl)propanamide

Triethylamine (75 mL, 54 mg, 0.54 mmol) was added to a mixture of 3-amino-2,3-dihydro-1-methyl-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-2-one (82 mg, 0.3 mmol), cyclohexanepropanoic acid (52 mL, 47 mg, 0.3 mmol), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (58 mg, 0.3 mmol) and 1-hydroxybenzotriazole (42 mg, 0.3 mmol) in DMF (1.5 mL). The mixture was stirred at room temperature for 18 h. and ethyl acetate (60 mL) was added. The mixture was washed with aqueous citric acid (10%), saturated aqueous sodium hydrogen carbonate and brine, dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with EtOAc/Hexane to give 3-cyclohexyl-N-(2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-thieno[2,3-e]-1,4-diazepin-3-yl)propanamide as a solid (56 mg, 46%).
m.p. 189-190°C.

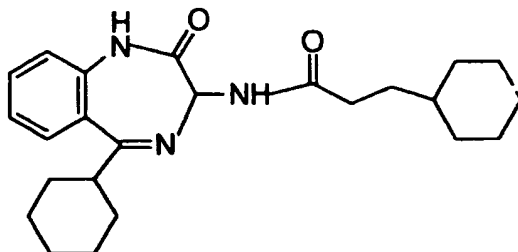
¹H (CDCl₃) 7.65-6.85 (8H, m), 5.65 (1H, d, J 8.0 Hz), 3.55 (3H, s), 2.40 (2H, t, J 7.0 Hz), and 1.80-0.85 (13H, m).

Anal. Calcd. for C₂₃H₂₇N₃O₂S.0.5H₂O:

C, 66.00; H, 6.74; N, 10.04.

Found: C, 66.25; H, 6.76; N, 9.83%.

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EXAMPLE 74**3-Cyclohexyl-N-(5-cyclohexyl-2,3-dihydro-2-oxo-1H-1,4-benzodiazepin-3-yl) propanamide**

Phenylmethyl N-[5-cyclohexyl-2,3-dihydro-2-oxo-1H-1,4-benzodiazepin-3-yl]carbamate (150 mg, 0.38 mmol) was dissolved in hydrogen bromide in acetic acid (30%, 0.5 mL). After 2 h., ether was added and the solid was collected and dried in vacuo. THF (3 mL) and triethylamine (0.45 mL, 33 mg, 0.32 mmol) were added and the mixture was stirred at room temperature for 3 h. In a separate flask, oxalyl chloride (38 mL, 56 mg, 0.44 mmol) was added to a solution of cyclohexanepropionic acid (61 mL, 56 mg, 0.36 mmol) and DMF (1 drop) in THF (2 mL) and the mixture was stirred at room temperature for 3 h. The two mixtures were combined, triethylamine (61 mL, 44 mg, 0.44 mmol) was added and the mixture was stirred at room temperature for 3 h. The solvent was evaporated under reduced pressure and ethyl acetate was added. The mixture was washed with water (2 x), saturated aqueous sodium hydrogen carbonate, water and brine, dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was recrystallized from i-PrOH to give 3-cyclohexyl-N-(5-cyclohexyl-2,3-dihydro-2-oxo-1H-1,4-benzodiazepin-3-yl)propanamide as a solid, m.p. 133-138°C.

dH (CDCl₃) 7.85 (1H, br s), 7.62-6.95 (5H, m), 5.40 (1H, d, J 8.7 Hz), 2.77 (1H, m), 2.34 (2H, m), and 2.05-0.75 (23H, m).

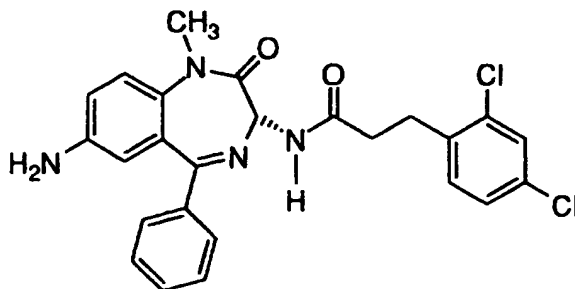
Anal. Calcd. for C₂₄H₃₃N₃O₂·0.7C₃H₇OH:

C, 71.64; H, 8.89; N, 9.60.

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Found: C, 71.28; H, 8.70; N, 9.82%.

EXAMPLE 75



(+)-N-[(3R)-7-Amino-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

Step A:

To a mixture of 3(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (*J. Org. Chem.* 1987, 52, 3232-3239) (3.98 g, 15.0 mmol) in concentrated sulfuric acid (15 mL) cooled in an ice-bath was added dropwise a solution of potassium nitrate (2.12 g, 21.0 mmol) in concentrated sulfuric acid (6 mL). The mixture was stirred with cooling for 2 h., then stirred at ambient temperature for 1.5 h. Ice (80 g) was added and the mixture was basified with concentrated ammonium hydroxide to pH 9. The resulting mixture was extracted with ethyl acetate (3 x 220 mL). The combined organic fractions were washed with brine, dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with chloroform/methanol (97:3). The material which eluted was further purified by flash column chromatography on silica gel, eluting with ethyl acetate/methanol (95:5). The material which eluted was stirred under *n*-butyl chloride (30 mL) and the solvent was evaporated under reduced pressure to give an inseparable mixture of 3(R)-amino-1,3-dihydro-1-methyl-7-nitro-5-phenyl-2H-1,4-benzodiazepin-2-one and 3(R)-amino-1,3-dihydro-1-

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methyl-7-nitro-5-(2-nitrophenyl)-2H-1,4-benzodiazepin-2-one (3.81 g) in a 3:1 ratio as a yellow solid.

δ H (CDCl₃) (mononitro compound) 8.43 (1H, dd, J 9, 3 Hz), 8.23 (1H, d, J 3 Hz), 7.59 (2H, m), 7.52 (2H,m), 7.44 (2H,m), 4.47 (1H,s), 3.53 (3H,s), and 2.42 (2H, br s); (dinitro compound) 8.49 (1H, dd, J 9, 3), 8.42 (1H, m), 8.18 (1H, d, J 3 Hz), 8.01 (1H, m), 7.67 (1H, t, J 6 Hz), 7.6-7.4 (2H, m), 4.52 (1H,s), 3.56 (3H,s), and 2.42 (2H, br s).

Step B:

A solution of 3-(2,4-dichlorophenyl)propionic acid (482 mg, 2.2 mmol), DMF (0.017 mL, 0.22 mmol), and thionyl chloride (0.24 mL, 3.3 mmol) in chloroform (2.5 mL) was heated at reflux for 1 h. The solvent was evaporated under reduced pressure to give 3-(2,4-dichlorophenyl)propionyl chloride (520 mg, 100%). To a solution of mixed 3(R)-amino-1,3-dihydro-1-methyl-7-nitro-5-phenyl-2H-1,4-benzodiazepin-2-one and 3(R)-amino-1,3-dihydro-1-methyl-7-nitro-5-(2-nitrophenyl)-2H-1,4-benzodiazepin-2-one (3:1) (621 mg, 2 mmol) and triethylamine (0.305 mL, 2.2 mmol) in methylene chloride (10 mL), was added a solution of 3-(2,4-dichlorophenyl)propionyl chloride (520 mg, 2.2 mmol) in methylene chloride (1.5 mL). The mixture was stirred for 30 min., the solvent was partially evaporated under reduced pressure, and the reaction mixture was purified by flash column chromatography on silica gel, eluting with methylene chloride/ether (90:10) to give a mixture of (+)-N-[(3R)-2,3-dihydro-1-methyl-7-nitro-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide and (+)-N-[(3R)-2,3-dihydro-1-methyl-7-nitro-2-oxo-5-(2-nitrophenyl)-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide (850 mg, 84%) in a 3:1 ratio as a solid white foam.

δ H (CDCl₃) (mononitro compound) 8.45 (1H, dd, J 9, 3 Hz), 8.25 (1H, d J 3 Hz), 7.54 (3H, m), 7.45 (2H, m), 7.38 (1H, d, J 2 Hz), 7.26-7.18 (4H, m), 5.50 (1H, d, J 8 Hz), 3.52 (3H, s), 3.10 (2H, m), and 2.70 (2H, m); (dinitro compound) 8.51 (1H, dd, J 9, 3 Hz), 8.40 (1H, m), 8.21 (1H, d J 3 Hz), 7.98 (1H, m), 7.68 (1H, t, J 6 Hz), 7.60 (1H, m), 7.44

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(1H, m), 7.26-7.15 (4H, m), 5.52 (1H, d, J 8 Hz), 3.55 (3H, s), 3.10 (2H, m), and 2.70 (2H, m).

Step C:

To a solution of mixed N-[(3R)-2,3-dihydro-1-methyl-7-nitro-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide and (+)-N-[(3R)-2,3-dihydro-1-methyl-7-nitro-2-oxo-5-(2-nitrophenyl)-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide (3:1) (770 mg, 1.5 mmol) in acetic acid (6 mL) was added dropwise in portions over 1.5 h. a solution of 15% titanium (III) chloride in 20-30% hydrochloric acid (7.8 mL, 9.0 mmol). The resulting solution was stirred 30 min., basified with 20% sodium hydroxide solution (pH 9), diluted with water (80 mL) and extracted with ethyl acetate (3 x 100 mL). The combined organic fractions were washed with brine, dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with ethyl acetate/hexane (75:25 increasing to 100:0). The first compound to elute was crystallized from ethyl acetate to give (+)-N-[(3R)-7-amino-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide (413 mg, 57%) as a pale yellow solid, m.p. 179-180°C, $[\alpha]_D^{+60.2}$ (c= 0.500, CHCl₃).

¹H NMR (CDCl₃) 7.60 (2H, d, J 7 Hz), 7.49-7.36 (5H, m) 7.24 (1H, d, J 9 Hz), 7.17 (2H, m), 6.99 (1H, dd, J 9, 3 Hz), 6.64 (1H, d, J 3 Hz), 5.54 (1H, d, J 8 Hz), 4.80-3.50 (2H, br s), 3.39 (3H, s), 3.09 (2H, t, J 8 Hz), and 2.68 (2H, dt, J_d 3, J_t 8 Hz).

Anal. Calcd. for C₂₅H₂₂Cl₂N₄O₂:

C, 62.38; H, 4.61; N, 11.64.

Found: C, 62.58; H, 4.68; N, 11.65%.

The second compound to elute was crystallized from ethyl acetate to give (+)-N-[(3R)-7-amino-2,3-dihydro-1-methyl-2-oxo-5-(2-aminophenyl)-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide (114 mg, 15%) as a pale yellow solid, m.p. 188-189°C,

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$[\alpha]_D +50.0^\circ$ ($c=0.100$, MeOH).

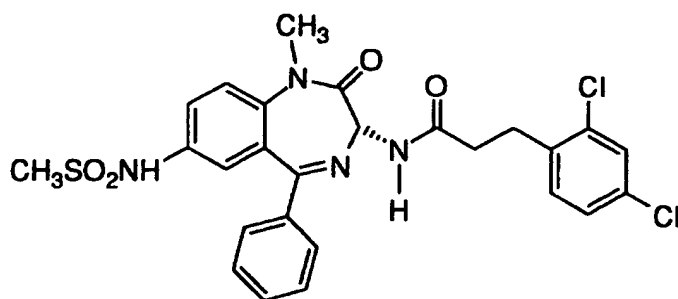
^1H (CDCl₃) 7.36 (2H, m), 7.25 (1H, d, J 9 Hz), 7.15 (3H, m), 7.00 (1H, m), 6.88 (2H, m), 6.79 (1H, m), 6.60 (1H, bs), 5.52 (1H, d, J 8 Hz), 4.10-2.80 (4H br s), 3.40 (3H, m), 3.09 (2H, t, J 8 Hz), and 2.69 (2H, m).

Anal. Calcd. for C₂₅H₂₃Cl₂N₅O₂·0.05EtOAc:

C, 60.43; H, 4.71; N, 13.99.

Found: C, 60.79; H, 4.74; N, 13.83%.

EXAMPLE 76



(+)-N-[(3R)-2,3-Dihydro-1-methyl-2-oxo-5-phenyl-7-methanesulfonamido-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

Methanesulfonyl chloride (0.040 mL, 0.52 mmol) was added to a solution of (+)-N-[(3R)-7-amino-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide (193 mg, 0.40 mmol) and pyridine (0.065 mL, 0.80 mmol) in methylene chloride (1.6 mL). The resulting solution was stirred 2 h. The solution was diluted with ethyl acetate (12 mL), washed with 1N HCl, water, saturated sodium bicarbonate solution, water, and brine (3 mL each), dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was dissolved in warm toluene, treated with charcoal, and filtered. The filtrate was diluted with hexane, the mixture was cooled, and the resulting precipitate was collected and

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dried in vacuo to give (+)-N-[(3R)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-7-methanesulfonamido-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide (152 mg, 68%) as a white solid, m.p. 130-148°C, $[\alpha]_D^{+111.6^\circ}$ ($c=0.500$, CHCl_3).

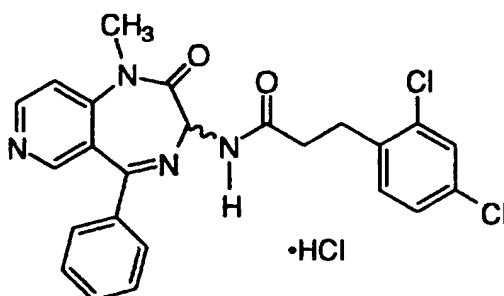
^1H (CDCl₃) 7.55-7.32 (9H, m), 7.24 (2H, dd, J 10, 2 Hz), 7.17 (1H, dd, J 9, 2 Hz), 7.05 (1H, d, J 3 Hz), 5.49 (1H, d, J 8 Hz), 3.41 (3H, s), 3.08 (2H, t, J 8 Hz), 2.97 (3H, s), and 2.71 (2H, dt, J_d 3, J_t 8 Hz).

Anal. Calcd. for C₂₆H₂₄Cl₂N₄O₄S:

C, 55.82; H, 4.32; N, 10.01.

Found: C, 56.12; H, 4.47; N, 9.89%.

EXAMPLE 77



N-(2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepin-3-yl)-3-(2,4-dichlorophenyl)propanamide hydrochloride

Step A:

To a solution of 2,3-dihydro-1-methyl-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepine-2-one (*J. Med. Chem.*, 1965, 8, 722-724) (1.63 g, 6.5 mmol) in toluene (32 mL) under argon cooled to -20°C (ice/methanol bath) was added potassium t-butoxide (1.83 g, 16.3 mmol). The resulting purple suspension was stirred 15 min. at -20°C and isoamyl nitrite (1.05 mL, 7.8 mmol) was added. The mixture was stirred at -20°C for 30 min., then poured into a mixture of water (50 mL), acetic acid (3 mL), and ethyl acetate (65 mL). The mixture was

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stirred to dissolve all solids and the layers were separated. The aqueous layer was extracted with ethyl acetate (65 mL). The combined organic fractions were washed with saturated sodium bicarbonate solution and brine (20 mL each), dried (Na_2SO_4), and the solvent was evaporated under reduced pressure. The residue was triturated with cold toluene and the solid was collected and dried in vacuo to give 2,3-dihydro-3-hydroxyimino-1-methyl-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepine-2-one (1.22 g, 67%) as a yellow solid, m.p. 223-224°C.

^1H NMR (CDCl_3) 8.92 (1H, bs), 8.73 (1H, d, J 7 Hz), 8.62 (1H, s), 7.80 (2H, dd, J 7, 1 Hz), 7.59 (1H, m), 7.48 (2H, m), 7.26 (1H, d, J 7 Hz), and 3.50 (3H, s).

Step B:

A mixture of 2,3-dihydro-3-hydroxyimino-1-methyl-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepine-2-one (1.77 g, 6.3 mmol) and freshly prepared Raney nickel (3.2 g) in 1:1 ethanol/methanol (190 mL) was shaken on a Parr hydrogenation apparatus under hydrogen (50 psi) for 4 h. The mixture was filtered through filter aid and the filtrate was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with methanol/chloroform/acetic acid (5:95:1 increasing to 10:90:1). The material which eluted was stirred under chloroform (30 mL) with potassium carbonate (0.3 g) and water (0.2 mL) for 5 min. The mixture was dried (Na_2SO_4) and the solvent was evaporated under reduced pressure to give 3-amino-2,3-dihydro-1-methyl-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepine-2-one (276 mg, 16%), as a yellow solid, m.p. 109-123°C. ^1H NMR (CDCl_3) 8.72 (1H, d, J 6 Hz), 8.58 (1H, s), 7.61 (2H, m), 7.51 (1H, m), 7.43 (2H, m), 7.26 (1H, m), 4.47 (1H, s), 3.50 (3H, s), and 2.1 (2H, bs).

High res. mass spectrum: Theoretical mass for $\text{C}_{15}\text{H}_{14}\text{N}_4\text{O}$ (M+1): 267.124586. Measured mass (M+1): 267.123654.

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Step C:

A solution of dicyclohexylcarbodiimide (87 mg, 0.42 mmol) in methylene chloride (0.17 mL) was added to a solution of 3-amino-2,3-dihydro-1-methyl-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepine-2-one (93 mg, 0.35 mmol) and 3-(2,4-dichlorophenyl)propionic acid (83 mg, 0.38 mmol) in tetrahydrofuran (0.5 mL) under argon. The resulting mixture was stirred for 5 h., filtered, and the filtrate was evaporated under reduced pressure. The residue was purified by preparative plate chromatography on silica gel eluting with methanol/chloroform/acetic acid (5:95:1). The purified material was stirred under chloroform (5 mL) with potassium carbonate (0.1 g) and water (2 drops) for 5 min. The mixture was dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was suspended in ethanol (2 mL) and ethanolic HCl (6.8 M, 0.147 mL) was added. The mixture was stirred, the resulting precipitate was collected and dried in vacuo to give N-(2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepin-3-yl)-3-(2,4-dichlorophenyl)propanamide hydrochloride (32 mg, 18%) as a white solid, m.p. 218-219°C.

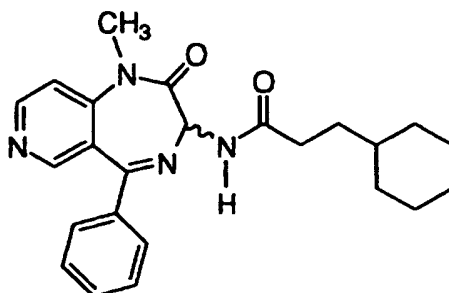
¹H (d₆-DMSO) 9.38 (1H, d, J 8 Hz), 8.86 (1H, bs), 8.59 (1H bs), 7.79 (1H, d, J 6 Hz), 7.56 (3H, m), 7.51 (2H, m), 7.39 (2H, m), 7.25 (1H, m), 7.16 (1H, m), 5.37 (1H, d, J 8 Hz), 3.44 (3H, s) 2.94 (2H, t, J 7 Hz), and 2.64 (2H, t, J 7 Hz).

Anal. Calcd. for C₂₄H₂₀Cl₂N₄O₂.HCl:

C, 57.22; H, 4.20; N, 11.12.

Found: C, 56.87; H, 4.18; N, 11.09%.

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EXAMPLE 78N-(2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepin-3-yl)-3-(cyclohexyl)propanamide

A solution of dicyclohexylcarbodiimide (87 mg, 0.42 mmol) in methylene chloride (0.17 mL) was added to a solution of 3-amino-2,3-dihydro-1-methyl-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepine-2-one (93 mg, 0.35 mmol) and cyclohexanepropionic acid (0.065 mL, 0.38 mmol) in tetrahydrofuran (0.5 mL) under argon. The resulting mixture was stirred for 5 h., filtered, and the filtrate was evaporated under reduced pressure. The residue was purified by preparative plate chromatography on silica gel eluting with methanol/chloroform/acetic acid (5:95:1). The purified material was stirred under chloroform (5 mL) with potassium carbonate (0.1 g) and water (2 drops) for 5 min. The mixture was dried (Na₂SO₄) and the solvent was evaporated under reduced pressure. The residue was crystallized from toluene to give N-(2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepin-3-yl)-3-(cyclohexyl)-propanamide (47 mg, 33%) as a white crystalline solid, m.p. 170-173°C.

¹H (CDCl₃) 8.75 (1H, d, J 6 Hz), 8.61 (1H, s), 7.58 (2H, m), 7.52 (1H, m), 7.45 (2H, m), 7.31 (1H, d, J 6 Hz), 7.21 (1H, d, J 8 Hz), 5.54 (1H, d, J 8 Hz), 3.51 (3H, s), 2.39 (2H, m), 1.73 (4H, m), 1.63 (3H, m), 1.85-1.12 (4H, m), and 0.94 (2H, m).

Anal. Calcd. for C₂₄H₂₈N₄O₂·0.10PhCH₃:

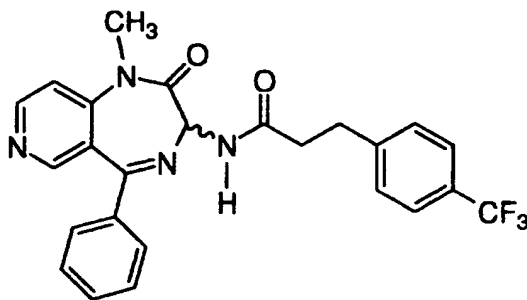
C, 71.70; H, 7.02; N, 13.54.

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Found: C, 71.78; H, 7.01; N, 13.57%.

Employing the procedure substantially as described above, but substituting 3-(4-trifluoromethylphenyl)-propionic acid for the cyclohexanepropionic acid, the following compound was prepared:

EXAMPLE 79



N-(2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-pyrido[4,3-e]-1,4-diazepin-3-yl)-3-(4-trifluoromethylphenyl)propanamide

m.p. 191-192°C.

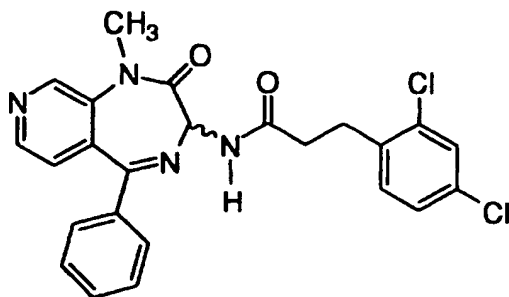
^1H (CDCl₃) 8.76 (1H, d, J 6 Hz), 8.61 (1H, s), 7.56 (4H, m), 7.52 (1H, m), 7.42 (2H, d, J 7 Hz), 7.38 (2H, m), 7.30 (1H, d, J 6 Hz), 7.22 (1H, d, J 8 Hz), 5.51 (1H, d, J 8 Hz), 3.50 (3H, s), 3.09 (2H, t, J 8 Hz), and 2.73 (2H, t, J 8 Hz).

Anal. Calcd. for C₂₅H₂₁F₃N₄O₂·0.20PhCH₃:

C, 65.39; H, 4.70; N, 11.56.

Found: C, 65.69; H, 4.64; N, 11.95%.

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EXAMPLE 80

N-(2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-pyrido[3,4-e]-1,4-diazepin-3-yl)-3-(2,4-dichlorophenyl)propanamide

Step A:

To a solution of 2,3-dihydro-1-methyl-5-phenyl-1H-pyrido[3,4-e]-1,4-diazepine-2-one (*Can. J. Chem.* 1987, 65, 1158-1161) (1.43 g, 5.7mmol) in toluene (28 mL) under argon cooled to -20°C (ice/methanol bath) was added potassium t-butoxide (1.59 g, 14.2 mmol). The resulting purple suspension was stirred 15 min. at -20 °C and isoamyl nitrite (0.92 mL, 6.8 mmol) was added. The mixture was stirred at -20°C for 30 min., then poured into a mixture of water (25 mL), acetic acid (2.5 mL), and ethyl acetate (55 mL). The mixture was stirred to dissolve all solids and the layers were separated. The aqueous layer was extracted with ethyl acetate (2 x 55 mL). The combined organic fractions were washed with saturated sodium bicarbonate solution and brine (20 mL each), dried (Na₂SO₄), and the solvent was evaporated under reduced pressure. The residue was triturated with hexane and the solid was collected and dried *in vacuo* to give 2,3-dihydro-3-hydroxyimino-1-methyl-5-phenyl-1H-pyrido[3,4-e]-1,4-diazepine-2-one (1.60 g, 100%) as a tan foam.

¹H (CDCl₃) 8.77 (1H, s), 8.50 (1H, d, J 4 Hz), 7.81 (2H, dd, J 8, 1 Hz), 7.60 (1H, m), 7.49 (3H, m), 7.32 (1H, d, J 5 Hz), and 3.55 (3H,s).

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Step B:

A solution of stannous chloride dihydrate (3.72 g, 16.5 mmol) in concentrated hydrochloric acid (11 mL) was added dropwise to 2,3-dihydro-3-hydroxyimino-1-methyl-5-phenyl-1H-pyrido[3,4-e]-1,4-diazepine-2-one (1.54 g, 5.5 mmol) cooled in an ice bath. The resulting solution was stirred at ambient temperature for 3 h. The solution was diluted with water (20 mL), basified with concentrated ammonium hydroxide (18 mL), and extracted with ether (4 x 75 mL). The combined organic fractions were washed with brine (30 mL), dried (Na_2SO_4), and the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography on silica gel, eluting with methanol/chloroform/acetic acid (5:95:1 increasing to 10:90:1). The material which eluted was stirred under chloroform (20 mL) with potassium carbonate (0.3 g) and water (2 drops) for 5 min. The mixture was dried (Na_2SO_4) and the solvent was evaporated under reduced pressure. The residue was stirred under hexane, and the resulting solid was collected to give 3-amino-2,3-dihydro-1-methyl-5-phenyl-1H-pyrido[3,4-e]-1,4-diazepine-2-one (241 mg, 16%) as a yellow solid, m.p. 94-118°C.

^1H (CDCl₃) 8.79 (1H, s), 8.48 (1H, d, J 5 Hz), 7.62 (2H, dd, J 8, 1 Hz), 7.51 (1H, m), 7.45 (2H, m), 7.24 (1H, dd, J 5, 1 Hz), 4.47 (1H, s), 3.55 (3H, s), and 2.2 (2H, bs).

Anal. Calcd. for $\text{C}_{15}\text{H}_{14}\text{N}_4\text{O} \cdot 0.25(\text{C}_2\text{H}_5)_2\text{O}$:

C, 67.46; H, 5.84; N, 19.67.

Found: C, 67.28; H, 5.66; N, 19.53%.

High res. mass spectrum: Theoretical mass for $\text{C}_{15}\text{H}_{14}\text{N}_4\text{O}$ (M+1):

267.124586. Measured mass (M+1): 267.123093.

Step C:

A solution of oxalyl chloride (0.023 mL, 0.26 mmol) in methylene chloride (0.2 mL) was added dropwise to a solution of 3-(2,4-dichlorophenyl)propionic acid (48 mg, 0.22 mmol) and DMF (1 drop) in methylene chloride (0.5 mL) cooled in an ice-bath. The resulting solution was stirred 1 h. with cooling. The solvent was

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evaporated under reduced pressure to give 3-(2,4-dichlorophenyl)-propionyl chloride (52 mg, 100%). To a solution of 3-amino-2,3-dihydro-1-methyl-5-phenyl-1H-pyrido[3,4-e]-1,4-diazepine-2-one (53 mg, 0.20 mmol) and pyridine (0.021 mL, 0.22 mmol) in methylene chloride (3 mL), was added a solution of 3-(2,4-dichlorophenyl)-propionyl chloride (52 mg, 0.22 mmol) in methylene chloride (0.5 mL). The mixture was stirred for 1 h., the solvent was partially evaporated under reduced pressure, and the reaction mixture was purified by flash column chromatography on silica gel, eluting with methanol/ether (5:95 increasing to 7.5:92.5). The material which eluted was crystallized from toluene/hexane to give N-(2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-pyrido[3,4-e]-1,4-diazepin-3-yl)-3-(2,4-dichlorophenyl)propanamide (38 mg, 38%) as a white crystalline solid, m.p. 220-221°C. δ H (CDCl₃) 8.81 (1H, s), 8.52 (1H, d, J 5 Hz), 7.56 (2H, dd, J 7, 2 Hz), 7.51 (1H, m), 7.44 (2H, d, J 6 Hz), 7.40 (1H, m), 7.27 (2H, m), 7.18 (2H, dd, J 8, 2 Hz), 5.48 (1H, d, J 8 Hz), 3.55 (3H, s), 3.10 (2H, t, J 7 Hz), and 2.71 (2H, dt, J_d 2 J_t 8 Hz).

Anal. Calcd. for C₂₄H₂₀Cl₂N₄O₂·0.25PhCH₃:

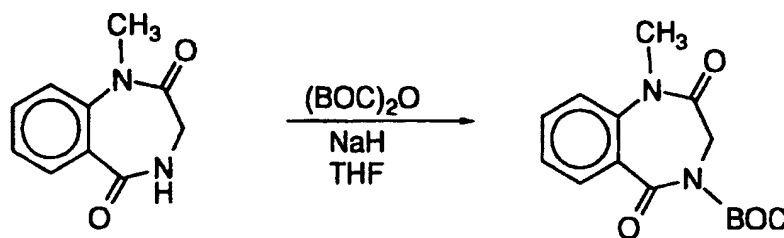
C, 63.06; H, 4.52; N, 11.43.

Found: C, 63.03; H, 4.48; N, 11.25%.

EXAMPLE 81

N-[2,3-Dihydro-1-methyl-2-oxo-5-isopropyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

Step A:

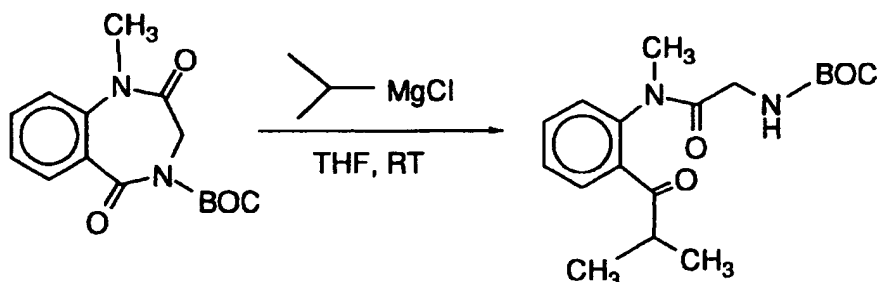


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To a solution of the benzodiazepine (1.0 g, 5.3 mmol) in THF (20 mL) at -78°C under argon was added 60% (NaH, 2.52 g, 6.3 mmol) Boc anhydride (1.27 g, 5.8 mmol) and the mixture stirred at -78°C for 1/2 hour. The reaction was then allowed to warm to 25°C and stirred for 2 hours before quenching into cold aq. NH_4Cl (10%) and extracting the product into ethyl acetate (3x50 mL). Concentration of the dried (Na_2SO_4) extracts gave an oil which was passed through silica (EtOAc/hexane) to give 1.35 g product (89%).

^1H NMR (CDCl_3) δ : 1.60 (s, 9H), 3.40 (s, 3H), 3.95 (brd, 1H), 4.80 (brd, 1H), 7.20 (d, 1H), 7.30 (q, 1H), 7.60 (t, 1H), 7.92 (d, 1H).

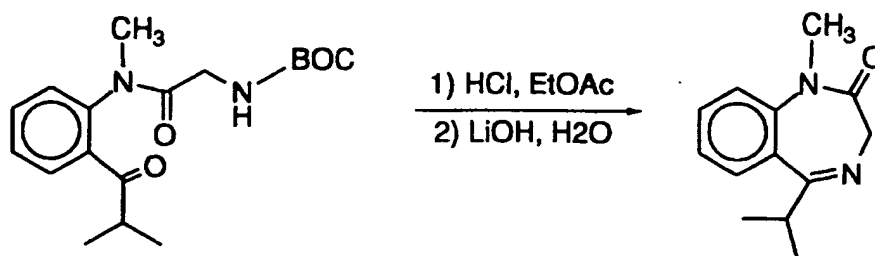
Step B:



To a solution of the BOC-benzodiazepine (4.0 g, 13.8 mmol) in THF (80 mL) under argon was rapidly added a solution of isopropylmagnesium chloride (2.0 M) in THF (7.66 mL, 15.3 mmol). The reaction was stirred for 1/2 hour, quenched into aq NH_4Cl (50 mL), and extracted with ethyl acetate (2x200 mL). The organic extracts were concentrated and chromatographed on silica (1:1, EtOAc/hexane) to give 1.55 g (34%) of product.

^1H NMR (CDCl_3) δ : 1.14 (d, 3H), 1.19 (d, 3H), 1.40 (s, 9H), 3.13 (s, 3H), 3.2-3.8 (m, 3H), 5.45 (brs, 1H), 7.28 (dt, 1H), 7.48 (dt, 1H), 7.56 (dt, 1H), 7.72 (dd, 1H).

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Step C:

To a 0°C solution of the isopropylphenone (1.55 g) in ethyl acetate was added anhydrous HCl gas over 90 min. The reaction was then concentrated in vacuo to give a solid which was dissolved in H₂O (40 mL) and the pH adjusted to 11.0 with 1N LiOH. After 30 min. at pH = 11.0 the pH was adjusted to 7.0 with 1N HCl and product extracted into ethyl acetate. The organic extracts were dried (Na₂SO₄), filtered and concentrated to give a solid 1.22 g, 100%.

¹H NMR (CDCl₃) δ: 0.95 (d, 3H), 1.30 (d, 3H), 3.16 (septet, 1H), 3.36 (s, 3H), 3.60 (d, 1H), 4.60 (d, 1H), 7.2-7.3 (m, 2H), 7.45-7.55 (m, 2H).

Step D:

The benzodiazepine obtained in Step C was converted to the oxime as described in Example 80 Step A.

Step E:

The oxime (2 gms) was dissolved in acetic acid (150 mL) and 10% Pd/C (1 gm) added. The mixture was stirred rapidly under an atmosphere of hydrogen for 90 min or until complete by HPLC. The reaction was filtered, the catalyst washed with methylene chloride (200 mL) and the filtrates concentrated in vacuo to an oil. The oil was dissolved in saturated aqueous sodium bicarbonate (100 mL) and product extracted with ethyl acetate (3 x 150 mLs). Concentration of the dried (Na₂SO₄) extracts gave 2.60 gms (97%).

Step F:

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The anine was coupled with 3-(2,4-dichlorophenyl)-propionic acid as described in Example 43 to yield N-(2,3-dihydro-1-methyl-2-oxo-5-isopropyl-1H-1,4-benzodiazepin-3-yl)-3-(2,4-dichlorophenyl)propanamide.

¹H NMR (CDCl₃) δ: 0.92 (d, 3H), 1.25 (d, 3H), 2.65 (dt, 2H), 3.05 (t, 2H), 3.15 (SepT, 1H), 3.40 (s, 3H), 5.38 (d, 1H), 7.0-7.6 (m, 8H).

The following compounds were prepared in a similar manner as described in Example 81, using the appropriate Grignard reagent in place of isopropyl magnesium chloride.

EXAMPLE 82

N-[2,3-dihydro-1-methyl-2-oxo-5-isopropyl-1H-1,4-benzodiazepin-3-yl]-3-cyclohexylpropanamide

m.p. 164-165°C

CHN: Anal. Calcd. for C₂₂H₃₁N₃O₂:

C, 71.51; H, 8.46; N, 11.37

Observed: C, 71.72; H, 8.39; N, 11.32

EXAMPLE 83

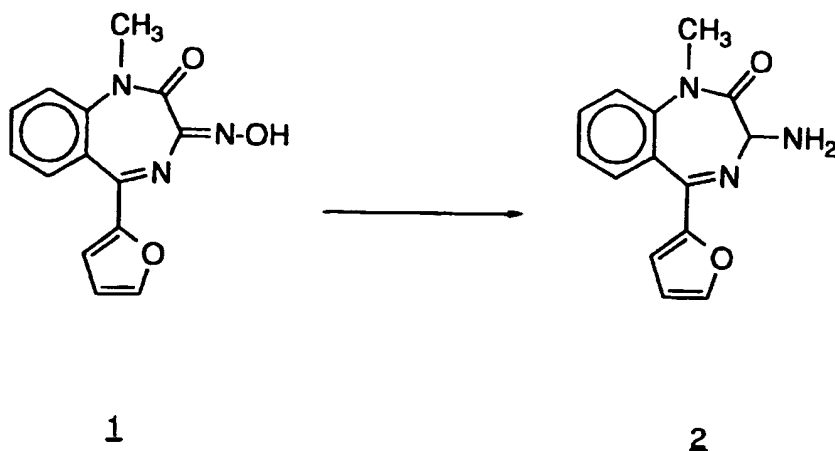
N-[2,3-dihydro-1-methyl-2-oxo-5-isopropyl-1H-1,4-benzodiazepin-3-yl]-3-(4-trifluoromethylphenyl)propanamide

m.p. 187-188°C

¹H NMR (CDCl₃) δ: 0.92 (d, 3H), 1.25 (d, 3H), 2.66 (dt, 2H), 3.04 (t, 2H), 3.15 (SepT, 1H), 3.40 (s, 3H), 5.38 (d, 1H), 7.14 (brd, 1H), 7.25-7.6 (m, 8H).

Employing substantially the same methods described in Example 80, but replacing Step E with the reduction method described below, the following compounds were prepared:

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To a solution of the oxime **1** (1.28 g, 0.0048 mole) in H₂O (130 ml) and THF (65 ml) was added sodium dithionite (Na₂S₂O₄) (13.0 g, 0.075 mole). The mixture was stirred for 2 hours then diluted with saturated aqueous sodium bicarbonate (50 ml) and product extracted into ethyl acetate (2 x 150 ml). The organic extracts were combined, dried over Na₂SO₄, filtered, and concentrated to give an oil (1.0 g). The oil was chromatographed on silica using ethyl acetate followed by 10% methanol/methylene chloride to give pure amine 0.778g (64%).

¹H NMR (DMSO) δ 3.32 (s, 3H), 4.30 (s, 1H), 6.64 (d, d, 1H), 6.76 (d, 1H), 7.35 (dt, 1H), 7.58-7.74 (m, 3H), 7.88 (m, 1H).

EXAMPLE 84

N-[2,3-dihydro-1-methyl-2-oxo-5-(2-furanyl)-1H-1,4-benzodiazepin-3-yl]-3-cyclohexylpropanamide

m.p. 168-169°C

CHN: Anal. Calcd. for C₂₃H₂₇N₃O₃:

C, 70.21; H, 6.92; N, 10.68

Observed: C, 70.15; H, 6.67; N, 10.64

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EXAMPLE 85

N-[2,3-dihydro-1-methyl-2-oxo-5-(2-furanyl)-1H-1,4-benzodiazepin-3-yl]-3-(4-trifluoromethylphenyl)propanamide

m.p. 155-157°C

CHN: Anal. Calcd. for C₂₄H₂₀N₃O₃F₃:

C, 63.29; H, 4.432; N, 9.23

Observed: C, 63.22; H, 4.44; N, 9.07

EXAMPLE 86

N-[2,3-dihydro-1-methyl-2-oxo-5-(2-furanyl)-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

m.p. 132-133°C

CHN: Anal. Calcd. for C₂₃H₁₉N₃O₃Cl₂

C, 60.54; H, 4.20; N, 9.21

Found: C, 60.62; H, 4.07; N, 9.07

EXAMPLE 87

N-[2,3-dihydro-1-methyl-2-oxo-5-(3-furanyl)-1H-1,4-benzodiazepin-3-yl]-3-cyclohexylpropanamide

m.p. 199-200°C

¹H NMR (CDCl₃) δ: 0.9-1.8 (brm, 3H), 2.38 (t, 2H), 3.42 (s, 3H), 5.55 (brd, 1H), 6.90 (s, 1H), 7.2-7.77 (m, 7H)

EXAMPLE 88

N-[2,3-Dihydro-1-methyl-2-oxo-5-(3-furanyl)-1H-1,4-benzodiazepin-3-yl]-3-(4-trifluoromethylphenyl)propanamide

m.p. 213-214°C

¹H NMR (CDCl₃) δ: 2.71 (dt, 2H), 3.05 (t, 2H), 3.42 (s, 3H), 5.72 (d, 1H), 6.82 (brs, 1H), 7.2-7.7 (m, 11H)

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EXAMPLE 89

N-[2,3-Dihydro-1-methyl-2-oxo-5-[2'-(4,4-dimethyl-2-oxazoliny)-phenyl]-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)-propanamide

The subject compound was prepared substantially as described in Example 81.

m.p. 194-195°C

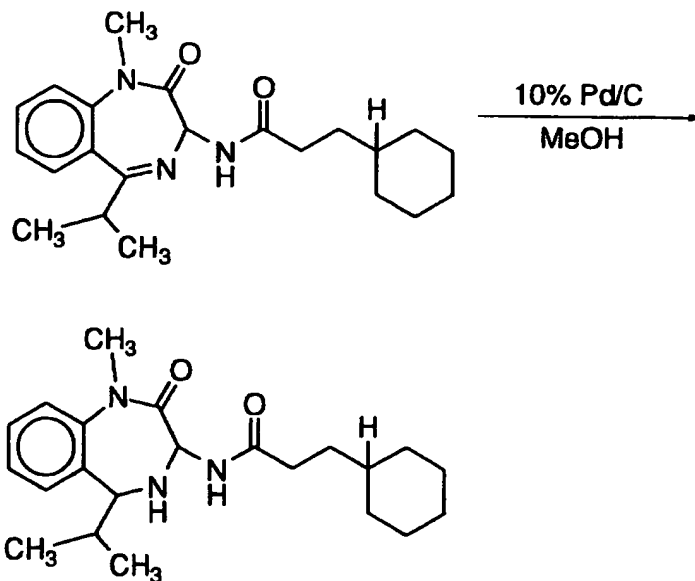
CHN: Anal. Calcd. for C₃₀H₂₈N₄O₃Cl₂

C, 63.95; H, 5.01; N, 9.94

Found: C, 63.70; H, 5.01; N, 9.96

EXAMPLE 90

N-[2,3,4,5-Tetrahydro-1-methyl-2-oxo-5-isopropyl-1H-1,4-benzodiazepin-3-yl]-3-cyclohexylpropanamide



A solution of N-[2,3-dihydro-1-methyl-2-oxo-5-isopropyl-

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1H-1,4-benzodiazepin-3-yl]-3-cyclohexylpropanamide (50 mg) in methanol (10 mL), containing 10% Pd/C (50 mg) was stirred under 1 atmosphere of hydrogen for 18 hours. Filtration of the reaction, concentration and crystallization from diethyl ether gave 21 mg N-[2,3,4,5-tetrahydro-1-methyl-2-oxo-5-isopropyl-1H-1,4-benzodiazepin-3-yl]-3-cyclohexylpropanamide.

CHN: Anal. Calcd. for C₂₂H₃₃N₃O₂

C, 71.12; H, 8.95; N, 11.31

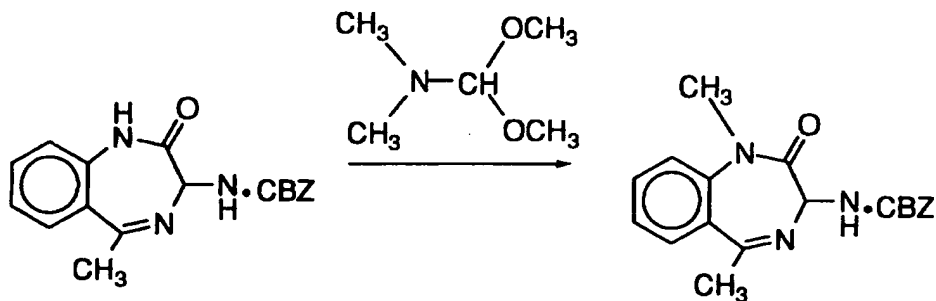
Observed: C, 70.98; H, 8.97; N, 11.15

m.p. 114-115°C

EXAMPLE 91

N-[2,3-dihydro-1-methyl-2-oxo-5-methyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

Step A:



To CBZ-benzodiazepine (250 mg, 0.776 mmol) in toluene (25 mL) at reflux was added dropwise a solution of DMF dimethylacetal (1.09 mL) in toluene (10 mL). The reaction was refluxed for 5 hours, cooled and concentrated to an oil. The oil was triturated with ether to give a white solid (124 mg).

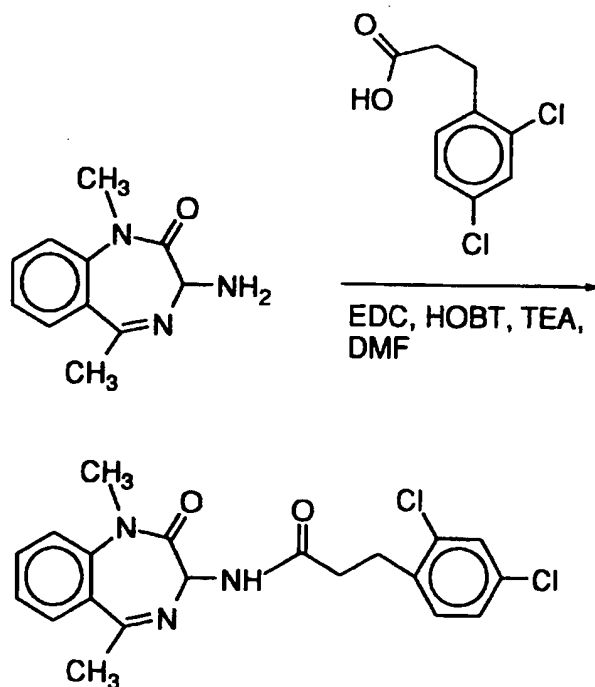
¹H NMR (CDCl₃) δ: 2.50 (s, 3H), 3.42 (s, 3H), 5.12-5.20 (m, 3H), 6.62 (d, 1H), 7.25-6.4 (m, 7H), 7.5-7.6 (m, 2H).

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Step B:

The CBZ-amine-N-methyl amide (190 mg) was treated with 30% HBr/AcOH (0.8 mL) for 1 hour at room temperature. The reaction mixture was poured into ether (10 mL) at 0°C and the solid filtered. Solid dissolved in 10% Aq. NaOH (5 mL) and CH₂Cl₂ (10 mL) and organic layer separated, dried (Na₂SO₄), filtered and concentrated to an oil (172 mg, 110%).

¹H NMR (CDCl₃) δ: 2.42 (s, 3H), 3.05 (brs, 2H), 3.40 (s, 3H), 4.40 (s, 1H), 7.2-7.6 (m, 4H).

Step C:

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N-[2,3-dihydro-1-methyl-2-oxo-5-methyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide was prepared in a similar manner as described previously in Example 43.

m.p. 194-195°C

CHN: Anal. Calcd. for $C_{20}H_{19}N_3O_2Cl_2$

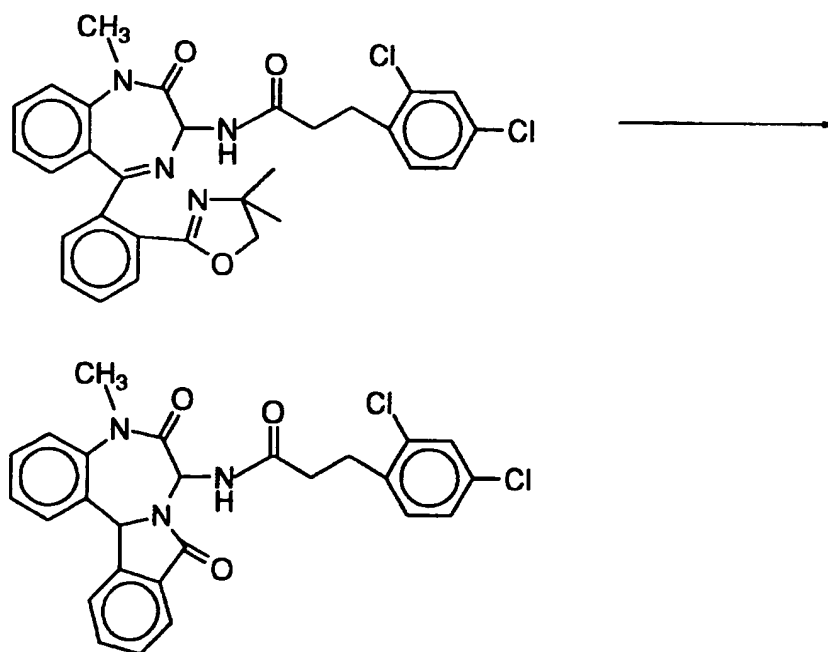
C, 59.42; H, 4.74; N, 10.39

Observed: C, 59.50; H, 4.74; N, 10.44

1H NMR ($CDCl_3$) δ : 2.49 (brs, 3H), 2.65 (dt, 2H), 3.05 (t, 2H), 3.42 (s, 3H), 5.35 (d, 1H), 7.1-7.6 (m, 8H).

EXAMPLE 92

N-[2,3-Dihydro-1-methyl-2-oxo-[4,5-a]-(1-oxo-1,3-dihydro-2H-isoindole)-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)-propanamide



To a solution of N-[2,3-dihydro-1-methyl-2-oxo-5-[2'-(4,4-dimethyl-2-oxazoliny)]phenyl]-1H-1,4-benzodiazepin-3-yl]-3-(2,4-

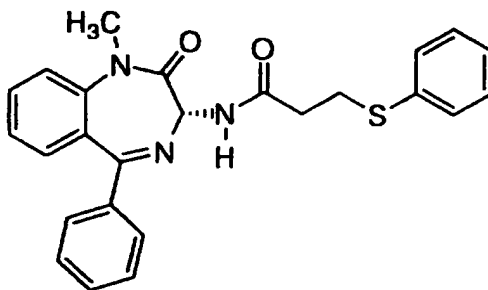
- 124 -

dichlorophenyl)propanamide (100 mg, 0.178 mmol) in methylene chloride was slowly added methyl trifluoromethanesulfonate (22 mL, 0.198 mmol). After stirring 5 minutes, sodium borohydride (7.6 mg, 0.20 mmol) in absolute ethanol (0.5 mL) was added and reaction stirred 30 min. the product was extracted into ethyl acetate and purified by column chromatography on silica (60% ethyl acetate/hexane) to give 30 mg N-[2,3-dihydro-1-methyl-2-oxo-[4,5-a]-(1-oxo-1,3-dihydro-2H-isoindole)-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)-propanamide.

^1H NMR (CDCl_3) δ : 2.70 (m, 2H), 3.12 (t, 2H), 3.55 (s, 3H), 5.68 (s, 1H), 5.90 (d, 1H), 6.85 (dd, 1H), 7.05 (brd, 1H), 7.1-7.5 (m, 9H), 7.85 (d, 1H).

MS $\text{M}^+ 494$.

EXAMPLE 93



3R-(+)-3-(Phenylthio)-N-[2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]propanamide

To a stirred solution of 3-bromopropionic acid (1.0g, 6.5mmol) in DMF (20 mL) was added K_2CO_3 (1.8 g, 13 mmol) and thiophenol (0.72 g, 6.5 mmol). This was heated to 50°C for 1h. The mixture was then diluted with 200 mL H_2O and extracted with 2 x 100 mL EtOAc. The combined organics were washed with 100 mL H_2O and dried with Na_2SO_4 . This was evaporated to give 1.52g of a colorless oil, 1.18g corrected for residual DMF by NMR.

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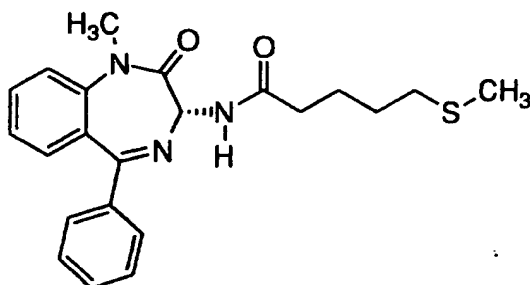
The above oil was taken up in 30 mL DMF and 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (2.45g, 12.8mmol) and 1-hydroxybenzotriazole hydrate (1.73g, 12.8mmol) were added. This was stirred for 5 min at rt. 3-(R)-Amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (0.66g, 2.6mmol) was then added and the reaction was stirred at rt overnight. The reaction was diluted with 200 mL H₂O and extracted with 2x150mL EtOAc. The combined organics were washed with 1x100mL H₂O, dried with Na₂SO₄ and evaporated. The residue was chromatographed over silica eluting with 2% MeOH:CHCl₃. Collected pure fractions, evaporated. Evaporated from diethyl ether to give 770mg of a white foam.

Anal. Calcd for C₂₅H₂₃N₃O₂S•0.05Hexane:

C, 70.04; H, 5.51; N, 9.69.

Found: C 69.91, H 5.40, N 9.78.

EXAMPLE 94



3R-(+)-5-(Methylthio)-N-[2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]propanamide

To an aqueous solution of K₂CO₃ (0.76g, 5.5mmol) was added 5-bromopentanoic acid and sodium thiomethoxide. This was stirred at rt overnight. The reaction was diluted with 50 mL H₂O and acidified to pH=0 with 6N HCl. Extracted with 2 x 50 mL EtOAc. Dried with Na₂SO₄, evaporated to give 0.55g of a yellow oil.

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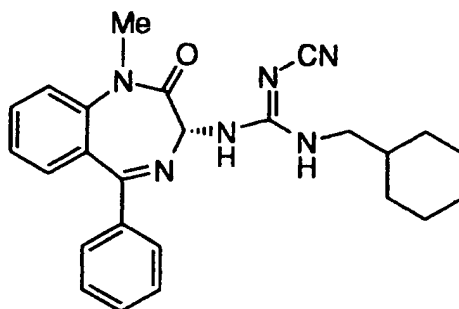
The above oil was taken up in 10 mL DMF and 1-(3-dimethyl-aminopropyl)-3-ethylcarbodiimide hydrochloride (1.30g, 6.8mmol) and 1-hydroxybenztriazole hydrate (0.92g, 6.8mmol) were added. 3-(R)-Amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (0.85g, 3.4 mmol) was then added and the reaction was stirred overnight at rt. The reaction was diluted with 100 mL H₂O and extracted with 2 x 50 mL EtOAc. Combined organics were dried with brine and Na₂SO₄, and evaporated to give yellow oil. The residue was chromatographed over silica eluting with 50:50 EtOAc:Hex to 100% EtOAc. Pure fractions were collected to give 1.33g of a colorless oil, 0.4g of which was chromatographed over silica eluting with 2% MeOH:CH₂Cl₂. Pure fractions were collected, and evaporated from ethyl ether:hexane to give a white powder mp. 61-65°C.

Anal. Calcd for C₂₂H₂₅N₃O₂S•0.35H₂O:

C, 65.76; H, 6.45; N, 10.46.

Found: C, 65.81; H, 6.21; N, 10.57.

EXAMPLE 95



N-cyano-N'-cyclohexylmethyl-N''-(1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl)guanidine

A solution of 3-(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (1g, 3.7 mmole) in acetonitrile (20 mL) was treated with diphenylcyanocarbonimidate (0.9 g, 3.7 mmole) and stirred at room temperature for thirty minutes. Cyclohexylmethyl-

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amine (0.84 g, 7.4 mmole) was then added and the reaction stirred at room temperature for two hours. The reaction was poured into 100 mL of 0.1 N HCl and extracted with 3 x 100 mL portions of ethyl acetate. The organic layers were combined and washed once with saturated sodium bicarbonate (50 mL), dried over anhydrous magnesium sulfate, filtered, and concentrated at reduced pressure. The residue was chromatographed on silica gel eluting with 50% ethyl acetate/hexane to give 0.875 g of the product. The analytical sample was crystallized from ethyl acetate.

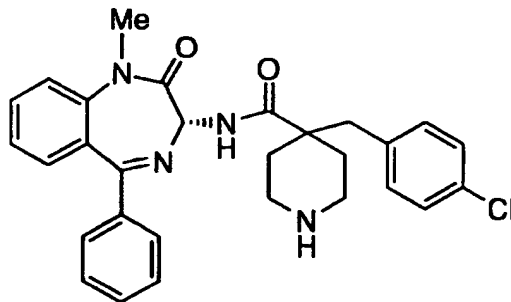
m.p. 158-161°C.

Anal. Calcd. for C₂₅H₂₈N₆O:

C, 70.07; H, 6.59; N, 19.61.

Found: C, 70.05; H, 6.59; N, 19.64%.

EXAMPLE 96



N-(1,3-Dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl)-4-(4-chlorobenzyl)-4-piperidinecarboxamide dihydrochloride

Step A: Preparation of N-tert-butyloxycarbonyl-4-(4-chlorobenzyl)-4-piperidinecarboxylic acid

A solution of N-Boc-ethylisonipecotate (51.4 g, 200 mmole) in THF (1L) at -60° C was treated with a solution of lithium bistrimethylsilyl amide (220 mL of a 1 N solution in THF, 220 mmole).

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After stirring at -60°C for 5 minutes, a solution of 4-chlorobenzyl chloride (33.8 g, 210 mmole) in THF (200 mL) was added and the reaction allowed to warm to room temperature. Most of the THF (about 800 mL) was removed by evaporation at reduced pressure. The remainder was poured into 1 L of 1 N HCl and extracted with two 800 mL portions of ethyl acetate. The organic layers were combined and washed once with saturated sodium bicarbonate (500 mL), dried over anhydrous magnesium sulfate, filtered, and concentrated at reduced pressure. The residue was chromatographed on silica gel eluting with 10%-20% ethyl acetate/hexane to give the product ester which was used directly. The material thus obtained was dissolved in THF (100 mL) and IPA (100 mL) and treated with 350 mL of 10 N NaOH. The mixture was heated to reflux for 30 hours. The reaction was cooled to room temperature and poured over a mixture of crushed ice (2 L), 6 N HCl (500 mL) and saturated potassium hydrogen sulfate (1 L). The mixture was extracted with two 1 L portions of ethyl acetate. The organic layers were combined and dried over anhydrous magnesium sulfate, filtered, and concentrated at reduced pressure to give 52 g of the product.

m.p. 179-180°C,

¹H NMR CDCl₃ δ 7.26 (d, J = 8 Hz, 2 H), 7.03 (d, J = 8 Hz, 2 H), 3.98 (m, 2H), 3.0-2.8 (m, 2H), 2.84 (s, 2H), 2.10-2.00 (m, 2H), 1.55-1.40 (m, 2H), 1.45 (s, 9H)

Step B: Preparation of N-(1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl)-4-(4-chlorobenzyl)-4-piperidinecarboxamide dihydrochloride

A mixture consisting of N-tert-butyloxycarbonyl-4-(4-chlorobenzyl)-4-piperidinecarboxylic acid (1.48 g, 4.18 mmole), 3-(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (1g, 3.7 mmole), hydroxybenzotriazole (1.17 g, 8.66 mmole), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (1.49 g, 7.70 mmole), diisopropylethyl amine (0.53 g, 4.13 mmole), and DMF (10 mL) was stirred at room temperature for 18 hours. The reaction was

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poured into 1 N HCl and extracted with ethyl acetate (4 X 50 mL). The organic layers were combined and washed once with saturated sodium bicarbonate (50 mL), once with saturated sodium chloride (50 mL), dried over anhydrous magnesium sulfate, filtered, and concentrated at reduced pressure. The residue was chromatographed on silica gel eluting with 25%-50% ethyl acetate/hexane to give 2.34 g of the product amide which was used directly. The material thus obtained was dissolved in ethyl acetate (50 mL) and HCl (g) was bubbled into the reaction for 5 minutes. The reaction was concentrated at reduced pressure and the residue recrystallized from ethyl acetate to give 1.13 g of the product as a pale yellow solid.

m.p. 190 - 195°C.

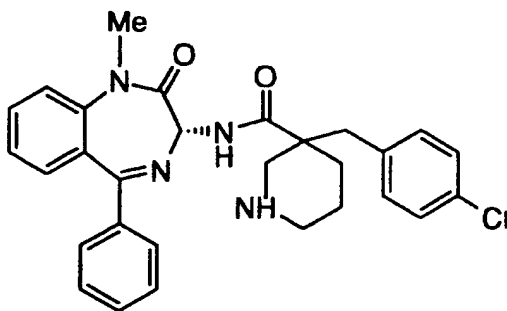
Anal. Calcd. for $C_{29}H_{29}ClN_4O_2 \cdot 2 HCl$:

C, 60.68; H, 5.44; N, 9.76.

Found: C, 60.47; H, 5.5; N, 9.42%.

Utilizing the procedures substantially as described above except substituting N-Boc-ethylnipecotate for N-Boc-ethyl isonipecotate there were obtained the following compounds

EXAMPLE 97



N-(1,3-dihydro-1-methyl-2-oxo-5-phenyl-2H-1,4-benzodiazepin-3-yl)-
3-(4-chlorobenzyl)-3-piperidinecarboxamide hydrochloride A + B
isomers

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Isomer A

m.p. 205 - 210°C.

Anal. Calcd. for $C_{29}H_{28}ClN_4O_2 \cdot HCl \cdot 0.5 CH_3CH_2OH \cdot 0.8 H_2O$:

C, 62.67; H, 6.07; N, 9.75.

Found: C, 62.69; H, 5.94; N, 9.42%.

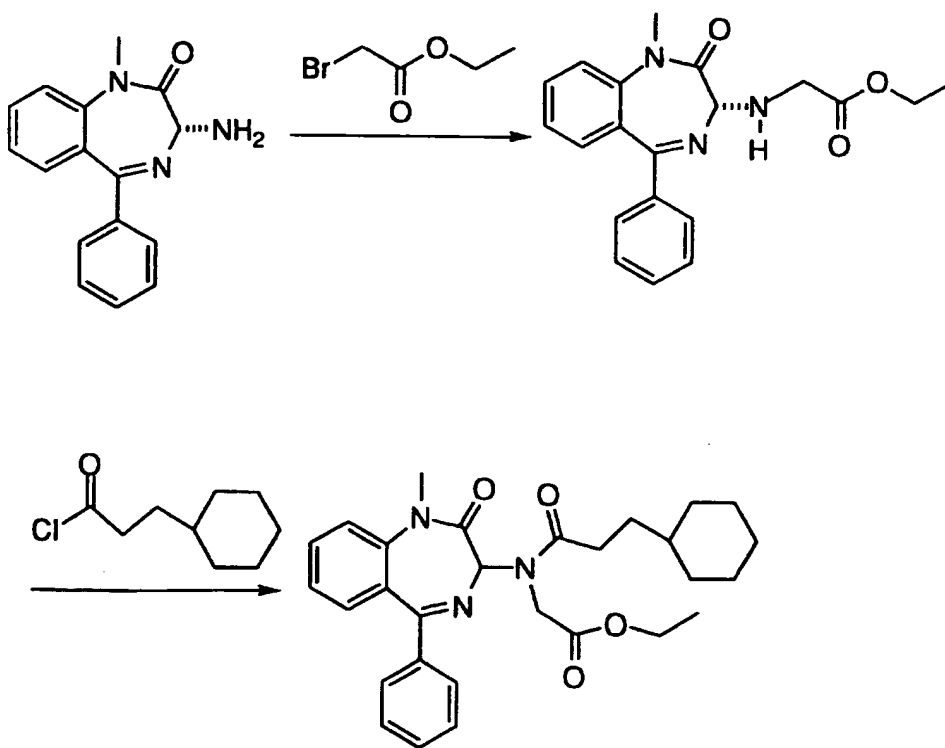
Isomer B

m.p. 200 - 205°C.

Anal. Calcd. for $C_{29}H_{28}ClN_4O_2 \cdot HCl \cdot 0.1 CH_3CH_2OCOCH_3 \cdot 1.6 H_2O$:

C, 61.39; H, 5.96; N, 9.74.

Found: C, 61.39; H, 5.66; N, 9.56%.

EXAMPLE 98

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(+)-3-Cyclohexyl-N-[2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-N-(ethoxycarbonylmethyl)propanamide

3-(R)-Amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (5.0 g, 18.8 mmol) in acetonitrile (100 mL) was mixed with ethyl bromoacetate (2.1 mL, 18.8 mmol) and sodium hydrogen carbonate (4.0 g) was suspended in the mixture. The mixture was stirred and heated at reflux for 2 h. After that time, the reaction was cooled to room temperature, diluted with 150 mL water, and extracted with ethyl acetate (3 x 100 mL). The organic layers were combined, dried with magnesium sulfate, gravity filtered, and the solvent was removed in vacuo. The resulting oil was chromatographed over silica in 3:1 ethyl acetate:hexane, yielding the mono-alkylated product (2.58 g, 39%) as well as the starting 1,4-benzodiazepin-2-one and bis-alkylated material. To a solution of 3-cyclohexylpropionic acid (1.0 g, 6.40 mmol) in methylene chloride (30 mL) was added oxalyl chloride (0.56 mL, 6.40 mmol) and catalytic (N,N)-dimethyl formamide (2 drops). After 0.5 h, a solution of the acetate (2.25 g, 6.40 mmol) in methylene chloride (10 mL) was added and the reaction was stirred for 0.25 h. The reaction was then diluted with methylene chloride (150 mL) and saturated aqueous sodium hydrogen carbonate (150 mL) was added. The aqueous portion was extracted again with methylene chloride (2 x 100 mL) and the organics were combined, dried with magnesium sulfate, gravity filtered, and the solvent was removed in vacuo. The resulting oil was chromatographed over silica with 1:1 ethyl acetate:hexane, yielding a foam that was crystallized with ether, giving 2.0 g (64%) of the product.

m.p. 120-122°C, $[\alpha]_D + 0.63^\circ$ (c=0.79; MeOH).

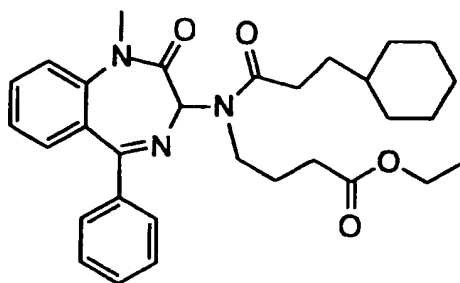
Anal. Calcd. for C₂₉H₃₅N₃O₄:

C, 71.14; H, 7.21; N, 8.58.

Found: C, 71.13; H, 7.13; N, 8.75%.

The following compound was prepared in a manner substantially as described above except substituting ethyl bromobutyrate for ethyl bromoacetate.

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EXAMPLE 99

3-Cyclohexyl-N-[2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-N-(ethoxycarbonylpropyl)propanamide

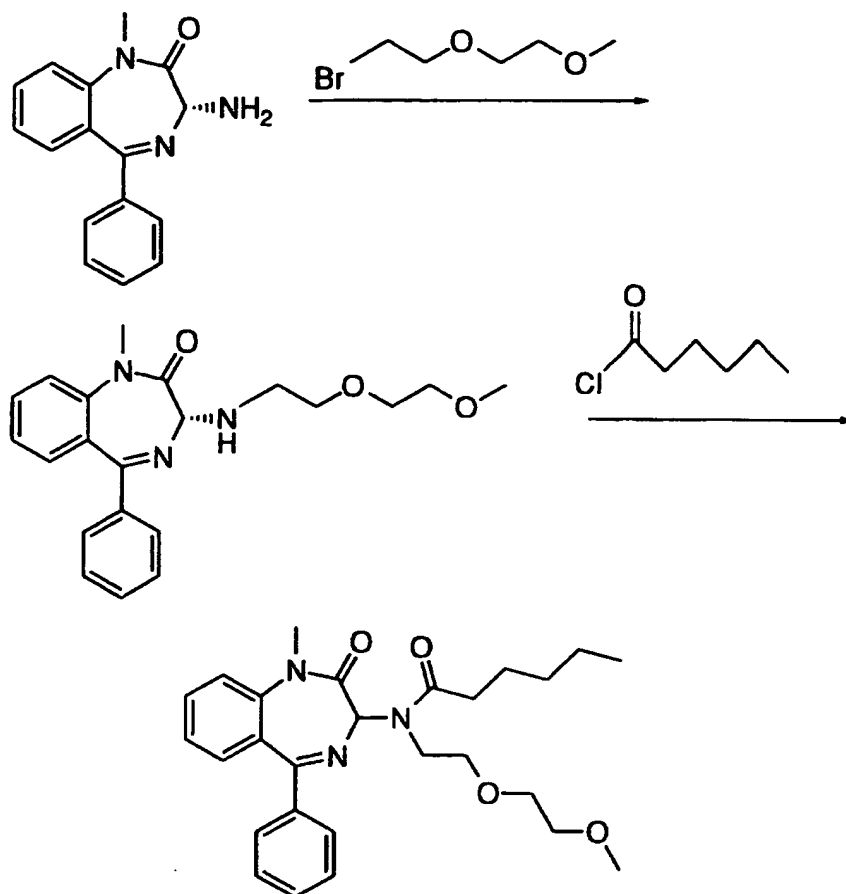
m.p. 103-105°C, $[\alpha]_D^{20}$ 0.00°; $c=0.85$; MeOH.

Anal. Calcd. for $C_{31}H_{39}N_3O_4 \cdot 0.40$ mol H_2O :

C, 70.94; H, 7.64; N, 8.01.

Found: C, 70.91; H, 7.44; N, 8.12%.

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EXAMPLE 100

N-[2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-
N-[2-(2-methoxyethoxy)ethyl]hexanamide

3-(R)-Amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (1.33 g, 5.0 mmol) in N,N-dimethyl formamide (30 mL) was mixed with 1-bromo-2-(2-methoxyethoxy)ethane (1.35 mL, 5.0 mmol) and triethylamine (1.0 mL). The mixture was stirred and heated at reflux for 4 h. After that time, the reaction was cooled to room temperature, diluted with 150 mL water, and extracted with ethyl acetate (3 x 100 mL). The organic layers were combined, dried with magnesium sulfate, gravity filtered, and the solvent was removed in

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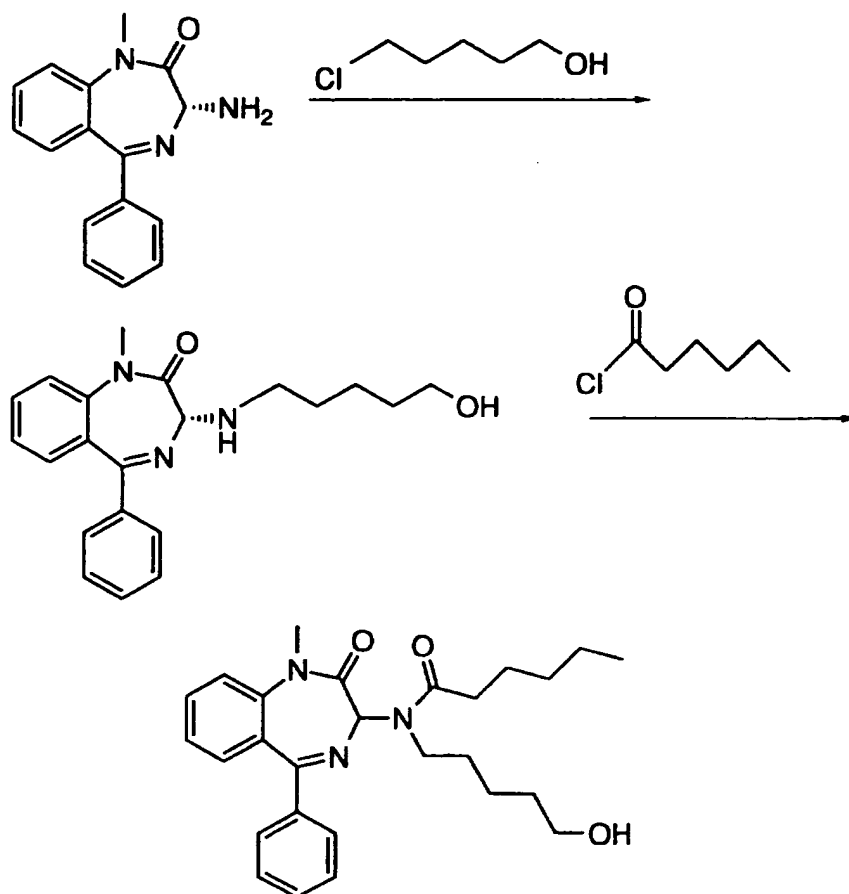
vacuo. The resulting oil was chromatographed over silica in 1:1 ethyl acetate:hexane, yielding the mono-alkylated product (1.2 g, 65%) as well as the starting 1,4-benzodiazepin-2-one and bis-alkylated material. To a solution of the mono-alkylated material (1.2 g, 3.27 mmol) in methylene chloride (20 mL) was added hexanoyl chloride (0.96 mL, 3.27 mmol) and the reaction was stirred for 0.25 h. The reaction was then diluted with methylene chloride (150 mL) and saturated aqueous sodium hydrogen carbonate (150 mL) was added. The aqueous portion was extracted again with methylene chloride (2 x 100 mL) and the organics were combined, dried with magnesium sulfate, gravity filtered, and the solvent was removed in vacuo. The resulting oil was chromatographed over silica with 1:1 ethyl acetate:hexane, yielding an oil, giving 580 mg (38%) of the product. $[\alpha]_D^{25} 0.00^\circ$; $c=0.27$; MeOH.

Anal. Calcd. for $C_{27}H_{35}N_3O_4 \cdot 0.80 \text{ mol H}_2\text{O}$:

C, 67.56; H, 7.69; N, 8.75.

Found: C, 67.56; H, 7.39; N, 8.85%.

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EXAMPLE 101

(+)-N-[2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-N-(5-hydroxypentyl)hexanamide

3-(R)-Amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (1.33 g, 5.0 mmol) in acetonitrile (40 mL) was mixed with 5-chloropentan-1-ol (0.61 g, 5.0 mmol) and sodium hydrogen carbonate (2.0 g) was suspended in the mixture. The mixture was stirred and heated at reflux for 12 h. After that time, the reaction was cooled to room temperature, diluted with 100 mL water, and extracted with ethyl acetate (3 x 75 mL). The organic layers were combined, dried with magnesium sulfate, gravity filtered, and the

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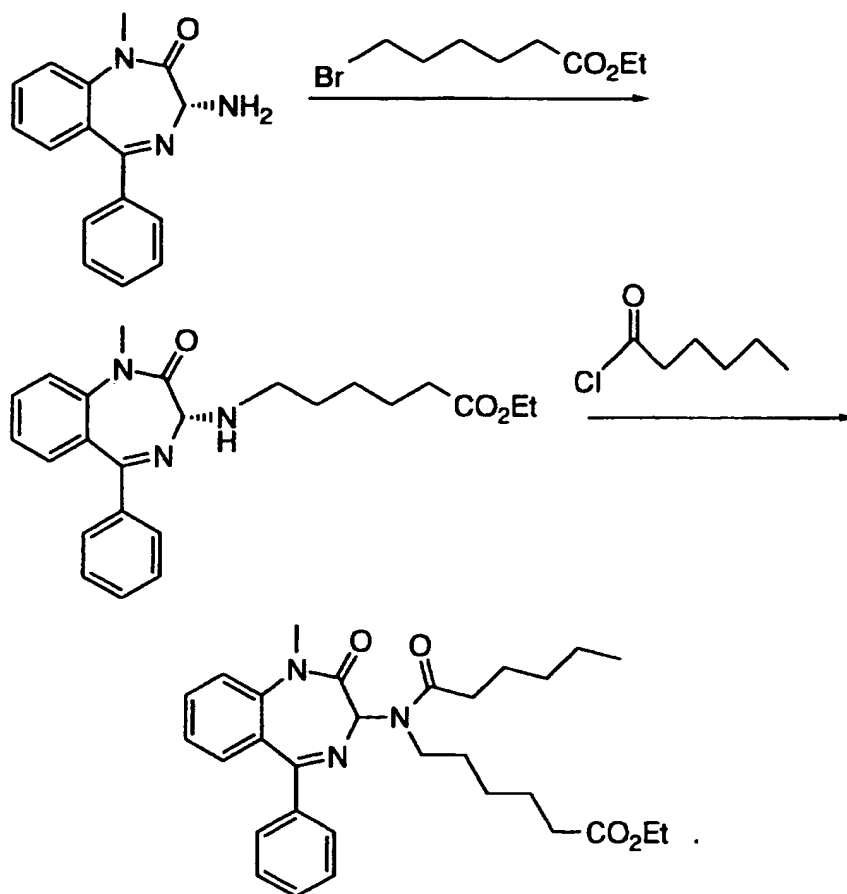
solvent was removed in vacuo. The resulting oil was chromatographed over silica in 1:49 methanol:chloroform yielding the mono-alkylated product (1.1 g, 62%) as well as the starting 1,4-benzodiazepin-2-one and bis-alkylated material. To a solution of the monoalkylated material (0.50 g, 1.42 mmol) in methylene chloride (30 mL) was added hexanoyl chloride (0.20 mL, 1.42 mmol) and the reaction was stirred for 0.25 h. The reaction was then diluted with methylene chloride (100 mL) and saturated aqueous sodium hydrogen carbonate (100 mL) was added. The aqueous portion was extracted with methylene chloride (2x75 mL) and the organics were combined, dried with magnesium sulfate, gravity filtered, and the solvent was removed in vacuo. The resulting oil was chromatographed over silica with 1:1 ethyl acetate:hexane, yielding a foam, giving 360 mg (64%) of the product. foam, $[\alpha]_D + 8.36^\circ$ (c=0.61, MeOH).

Anal. Calcd. for $C_{27}H_{35}N_3O_2 \cdot 0.25 \text{ mol } H_2O$:

C, 71.42; H, 7.88; N, 9.25.

Found: C, 71.47; H, 7.89; N, 9.12%.

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EXAMPLE 102

(+)-N-[2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-N-(ethoxycarbonylpentyl)hexanamide

3-(R)-Amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (1.33 g, 5.0 mmol) in acetonitrile (40 mL) was mixed with ethyl 6-bromohexanoate (0.89 mL, 5.0 mmol) and sodium hydrogen carbonate (2.0 g) was suspended in the mixture. The mixture was stirred and heated at reflux for 10 h. After that time, the reaction was cooled to room temperature, diluted with 100 mL water, and extracted with ethyl acetate (3x75 mL). The organic layers were combined, dried with magnesium sulfate, gravity filtered, and the

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solvent was removed in vacuo. The resulting oil was chromatographed in 1:49 methanol:chloroform, yielding the mono-alkylated product (0.56 g, 28%) as well as the starting 1,4-benzodiazepin-2-one and bis-alkylated material. To a solution of the mono-alkylated material (0.56 g, 1.37 mmol) in methylene chloride (20 mL) was added hexanoyl chloride (0.19 mL, 1.37 mmol) and the reaction was stirred for 0.25 h. The reaction was then diluted with methylene chloride (100 mL) and saturated aqueous sodium hydrogen carbonate (100 mL) was added. The aqueous portion was extracted again with methylene chloride (2x75 mL) and the organics were combined, dried with magnesium sulfate, gravity filtered, and the solvent was removed in vacuo. The resulting oil was chromatographed over silica with 1:1 ethyl acetate:hexane, yielding a foam, giving 0.40 g (58%) of the product.
m.p. 59-65°C, $[\alpha]_D^{25}$ (+)52.7° (c=0.48, MeOH).

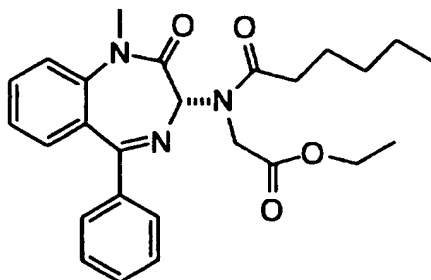
Anal. Calcd. for $C_{30}H_{39}N_3O_4 \cdot 0.20 \text{ mol } CH_2Cl_2$:

C, 69.4; H, 7.6; N, 8.04.

Found: C, 69.44; H, 7.68; N, 7.71%.

The following compound was prepared in a manner substantially as described above except substituting ethyl bromoacetate for ethyl 6-bromohexanoate.

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EXAMPLE 103

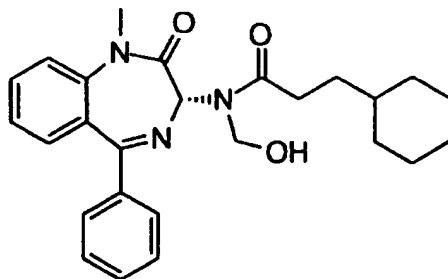
(+)-N-[2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-N-(ethoxycarbonylmethyl)hexanamide

foam, $[\alpha]_D + 2.04^\circ$ (c=0.98; MeOH).

Anal. Calcd. for $C_{26}H_{31}N_3O_4$:

C, 69.47; H, 6.95; N, 9.35.

Found: C, 69.41; H, 7.03; N, 9.26%.

EXAMPLE 104

(+)-3-Cyclohexyl-N-[2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-N-(hydroxymethyl)propanamide

(+)-3-Cyclohexyl-N-[2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]propanamide (2.0 g, 5.0 mmol) was dissolved in tetrahydrofuran (30 mL), cooled to 0°C and methyl

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magnesium chloride (3M, 2.0 mL) was added. After 0.25 h, paraformaldehyde (0.15 g, 10 mmol) was added, and the mixture was allowed to warm to room temperature. The reaction was then diluted with ethyl acetate (150 mL) and saturated aqueous sodium hydrogen carbonate (150 mL) was added. The aqueous portion was extracted again with ethyl acetate (2 x 100 mL) and the organics were combined, dried with magnesium sulfate, gravity filtered, and the solvent was removed *in vacuo*. The resulting oil was chromatographed over silica with 1:1 ethyl acetate:hexane, yielding a foam (0.80 g, 37%). foam, $[\alpha]_D + 124^\circ$ ($c=0.69$, MeOH).

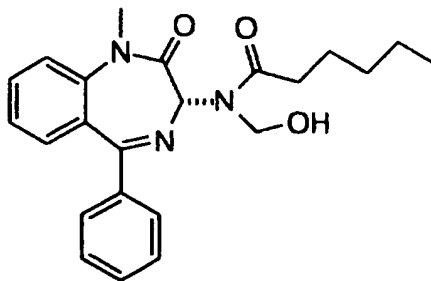
Anal. Calcd. for $C_{26}H_{31}N_3O_3$:

C, 72.03; H, 7.21; N, 9.69.

Found: C, 71.66; H, 7.08; N, 9.78%.

The following compound was prepared in a manner substantially as described above starting from (+)-N-[2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]hexanamide.

EXAMPLE 105



(+)-N-[2,3-Dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-N-(hydroxymethyl)hexanamide

m.p. 154-156°C, $[\alpha]_D + 190.8^\circ$ ($c=0.24$, MeOH).

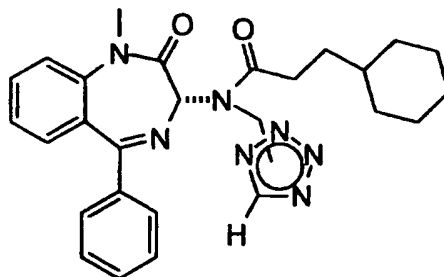
Anal. Calcd. for $C_{23}H_{27}N_3O_3 \cdot 0.30$ mol H_2O :

C, 69.26; H, 6.97; N, 10.53.

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Found: C, 69.29; H, 6.81; N, 10.6%.

EXAMPLE 106



(+)-3-Cyclohexyl-N-[2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-N-(tetrazolylmethyl)propanamide

(+)-3-Cyclohexyl-N-[2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepin-3-yl]-N-(hydroxymethyl)propanamide (0.67 g, 1.56 mmol) was dissolved in methylene chloride (100 mL), along with tetrazole (0.33 g, 4.7 mmol), and then N,N-diisopropyl-dibenzyl-phosphoramidite (1.07 g, 3.1 mmol). After 2 h, the mixture was diluted with methylene chloride (150 mL), and extracted with saturated aqueous sodium hydrogen carbonate (3 x 100 mL). The organic layers were combined, dried with magnesium sulfate, gravity filtered, and the solvent was removed in vacuo. The resulting oil was chromatographed twice over silica with 1:1 ethyl acetate:hexane, yielding two constitutional isomers, a (65 mg, 9%) and b (56 mg, 7.5%).

Isomer A:

m.p. 96-98°C, $[\alpha]_D^{25} +188.9^\circ$ (c=0.19, MeOH).

Anal. Calcd. for C₂₇H₃₁N₇O₂·0.30 mol TFA:

C, 63.78; H, 6.07; N, 18.86.

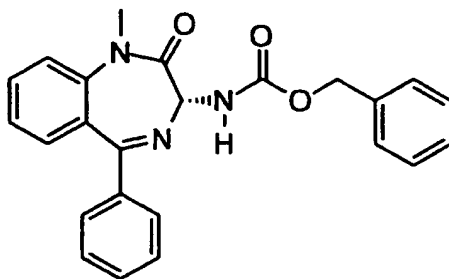
Found: C, 63.7; H, 6.12; N, 18.76%.

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Isomer B:m.p. 92-95°C, $[\alpha]_D^{+81.3^\circ}$ (c=0.31, MeOH).Anal. Calcd. for $C_{27}H_{31}N_7O_{20.35}$ mol TFA:

C, 63.31; H, 6.01; N, 18.66.

Found: C, 63.35; H, 6.02; N, 18.74%.

EXAMPLE 107**3R-(+)-3-(Benzyloxycarbonylamino)-2,3-dihydro-1-methyl-2-oxo-5-phenyl-1H-1,4-benzodiazepine**

To a stirring solution of 3-(R)-amino-1,3-dihydro-1-methyl-5-phenyl-2H-1,4-benzodiazepin-2-one (2.0 g, 7.5 mmol) in methylene chloride (45 mL) at 0°C was added benzyl chloroformate (1.2 mL, 8.3 mmol) and the reaction was allowed to warm to room temperature. The reaction mixture was diluted with methylene chloride (150 mL), and extracted with saturated aqueous sodium hydrogen carbonate (150 mL). The aqueous portion was extracted with methylene chloride (2 x 100 mL) and the organics were combined, dried with magnesium sulfate, gravity filtered, and the solvent was removed in vacuo. The resulting oil was chromatographed over silica with 1:1 ethyl acetate:hexane, yielding a white foam (3.0 g, 99.7%) $[\alpha]_D^{+57.5^\circ}$ (c=1.17; MeOH).

Anal. Calcd. for $C_{24}H_{20}N_3O_3 \cdot 0.70$ mol $H_2O \cdot 0.15$ mol $CHCl_3$:

C, 67.62; H, 5.06; N, 9.8.

Found: C, 67.6; H, 5.02; N, 9.75%.

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The following compounds were prepared substantially as described in Example 81.

EXAMPLE 108

N-[2,3-Dihydro-1-methyl-2-oxo-5-ethyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

m.p. 156-158°C.

CHN: Anal. Calcd. for $C_{21}H_{21}Cl_2N_3O_2 \cdot 0.5 H_2O$:

C, 59.02; H, 5.19; N, 9.83.

Found: C, 58.99; H, 4.89; N, 9.88.

EXAMPLE 109

N-[2,3-Dihydro-1-methyl-2-oxo-5-t-butyl-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

m.p. 170-171°C.

CHN: Anal. Calcd. for $C_{23}H_{25}Cl_2N_3O_2 \cdot 0.7 H_2O$:

C, 60.18; H, 5.80; N, 9.16.

Found: C, 60.17; H, 5.30; N, 9.30.

EXAMPLE 110

N-[2,3-Dihydro-1-methyl-2-oxo[4'-(4,4-dimethyl-2-oxazoliny)]phenyl]-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

m.p. 188-190°C.

CHN: Anal. Calcd. for $C_{30}H_{28}N_4O_3Cl_2$:

C, 63.95; H, 5.01; N, 9.94.

Found: C, 63.96; H, 5.02; N, 10.08.

EXAMPLE 111

N-[2,3-Dihydro-1-methyl-2-oxo-5-(4-methoxyphenyl)-1H-1,4-benzodiazepin-3-yl]-3-(2,4-dichlorophenyl)propanamide

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m.p. 188-189°C.

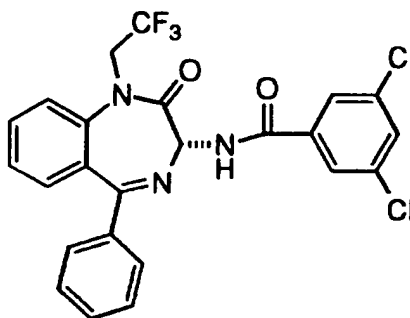
CHN: Anal. Calcd. for $C_{26}H_{23}Cl_2N_3O_3 \cdot 0.45 H_2O$:

C, 62.91; H, 4.67; N, 8.47.

Found: C, 61.89; H, 4.78; N, 8.33.

EXAMPLE 112

(+)-3,5-Dichloro-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]benzamide.



Step A: Preparation of 2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepine.

A solution of 5-phenyl-1,4-benzodiazepine-2-one (*J. Org. Chem.*, 1962, 27, 3788) (50 g, 0.211 mole) in DMF (100 mL) was treated with cesium carbonate (103.5 g, 0.317 mole) and trifluoroethyl iodide (109.7 g, 0.525 mole). The mixture was stirred at 50°C overnight. The reaction mixture was then poured into water (2 L) and extracted with ethyl acetate (3 X 1 L). The combined ethyl acetate fractions were dried over anhydrous magnesium sulfate, filtered and concentrated at reduced pressure. The residue was crystallized from ethyl ether to give 45 g (68 %) of the product. MP = 130 - 131°C; 1H NMR ($CDCl_3$, 300 MHz) δ 7.65-7.60 (m, 2H), 7.60-7.45 (m, 5H), 7.40-7.20 (m, 2H), 5.25 (dq, J = 14, 8.6 Hz, 1H), 4.82 (d, J = 10.5 Hz, 1H), 4.15 (app sextet, J = 8.6 Hz, 1H), 3.81 (d, J = 10.5 Hz, 1H).

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Step B: Preparation of 3-Azido-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepine.

To a stirring solution of 5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepine (70 g, 0.22 mol) in THF (1500 mL) cooled to -70°C was added potassium tert-butoxide (1.1 eq, 0.24 mol, 240 mL of a 1 N solution in THF) dropwise over 15 min. A solution of 2,4,6-triisopropylbenzenesulfonylazide (74.8 g, 0.24 mol) in THF (250 mL) was added over 5 min. This was stirred for 10 minutes and acetic acid (40 mL, 0.63 mol) was added and the reaction allowed to warm to ambient temperature. The reaction was poured into satd. NaHCO₃ (1500 mL) and ethyl acetate (1L). The phases were separated and the aqueous phase was extracted with ethyl acetate (500 mL). The combined organic layers were washed with water (500 mL) then brine (300 mL). The organic layers were dried with Na₂SO₄ and evaporated to a brown foam. This was triturated with ethyl ether to give 65 g of a white powder. The filtrate was concentrated and chromatographed over silica gel eluting with 30% ethyl acetate/hexane to give another 8.9 g. The combined yield was 74 g (93%). MP = 159 - 160°C; ¹H NMR (CDCl₃, 300 MHz) δ 7.70-7.26 (m, 9H), 5.28-5.12 (m, 1H), 4.63 (s, 1H), 4.35-4.10 (m, 1H).

Step C: Preparation of racemic 3-Amino-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepine.

To a stirring solution of 3-Azido-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-2,3-dihydro-1H-benzo[e][1,4]diazepine (83.4 mmol, 30 g) in 300 mL ethanol and 150 mL THF was added 10% Pd/C (10 wt%, 3.0 g). Hydrogen gas was bubbled through the solution for 8 h. The reaction was filtered and evaporated under reduced pressure. The residue was crystallized from ethyl ether to give 20.0 g of white crystals. Another 4 g was recovered from evaporation and recrystallization of the filtrates. Combined yield, 86.7%.

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MP = 141 - 143°C;
¹H NMR (CDCl₃, 300 MHz) δ 7.70-7.26 (m, 9H), 5.28-5.12 (m, 1H),
4.57 (s, 1H), 4.35-4.10 (m, 1H).

Step D: Preparation of 2-Amino-N-[2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl]-3-phenylpropionamide

To a stirring solution of 3-Amino-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-2,3-dihydro-1H-benzo[e][1,4]diazepine (92.2 mmol, 30.74g) in DMF (300mL) was added N-Benzyloxy-D-Phenylalanine (92.2 mmol, 27.6g), 1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.12mol, 22.95g) and 1-hydroxybenzotriazole hydrate (46.1mmol, 6.23g). This was stirred at room temperature for 2h. The reaction was then diluted with 1L of 10% KHSO₄ and extracted with ethyl acetate (2x600 mL). The organic layers were combined and washed with saturated sodium hydrogen carbonate (600mL). They were dried with brine and sodium sulfate and evaporated under reduced pressure. 66.58g of an orange foam, which contained ethyl acetate by NMR. NMR ¹H (CDCl₃) δ 7.75-7.18 (m, 20H), 5.62-5.55 (m, 1H), 5.48-5.00 (m, 4H), 4.72-4.60 (m, 1H), 4.25-4.05 (m, 1H) 3.32-3.05 (m, 2H). This material was carried on without further purification.

To a stirring solution of 2-(N-Benzyloxyamino)-N-[2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl]-3-phenyl propionamide in 1L ethanol was added 10% Pd/C (15 wt%) and hydrogen was bubbled through the reaction for 2h and then left stirring under 1 atm. hydrogen overnight. Hydrogen was bubbled through the reaction for an additional three hours the following morning. The reaction was then filtered, the catalyst was rinsed with 1L methylene chloride and evaporated under reduced pressure. The resulting solid was dried under vacuum overnight to give 44.46g of a white solid. This was chromatographed over silica, eluting with 1% MeOH:EtOAc. The pure upper R_f fractions were collected and

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evaporated under reduced pressure. The mixed fractions were collected, evaporated and rechromatographed. The pure fractions were collected and combined with the above pure fractions to get a combined yield of 18.11g, 83.5% of the upper R_f diastereomer. ¹H NMR (CDCl₃, 300 MHz) δ 8.94 (d, J=8.6Hz, 1H), 7.65-7.10 (m, 9H), 5.64 (d, J=8.6 Hz, 1H), 5.28-5.12 (m, 1H), 4.57 (s, 1H), 4.35-4.10 (m, 1H), 3.71 (dd, J=9.8 and 3.9 Hz, 1H), 3.34 (dd, J=13.9 and 3.9 Hz, 1H), 2.79 (dd, J=13.9 and 10.0 Hz, 1H). The Absolute stereochemistry at C-3 of the benzodiazepine ring was determined to be (R) by X-Ray analysis.

The lower R_f material corresponding to C-3(S) was isolated as well.

Step E: Preparation of 3(R)-(+)-3-Amino-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4] diazepine.

To a stirring solution of 2-Amino-N-[2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-2,3-dihydro-1H-benzo[e][1,4]diazepin-3-yl]-3-phenylpropionamide (13.6 g, 28.3 mmol) in methylene chloride (136 mL) was added phenyl isothiocynate (3.87 mL, 34.0 mmol). This was stirred overnight at ambient temperature. The reaction was then cooled in ice, trifluoroacetic acid (2.73 mL, 0.283 mol) added and the reaction allowed to warm to ambient temperature. After stirring at ambient temperature for 2.5 hours the reaction was evaporated under reduced pressure, chromatographed with 90:10:1:1 methylene chloride: methanol:acetic acid:water. The low R_f spot was collected and evaporated under reduced pressure with no heat. The residue was taken up in 600 mL methylene chloride and washed with 300 mL saturated NaHCO₃ and 300 mL water. The solution was dried over Na₂SO₄ and evaporated under reduced pressure. The residue was crystallized from ethyl acetate:hexanes to give 6.65 g of a white powder.

MP = 162 - 164°C;

¹H NMR (CDCl₃, 300 MHz) δ 7.70-7.26 (m, 9H), 5.28-5.12 (m, 1H), 4.57 (s, 1H), 4.35-4.10 (m, 1H).

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 $[\alpha]_D = +72.9^\circ$ ($c=0.7$, MeOH)

The (-)-3S enantiomer was prepared in the same fashion from the Lower Rf product of Step D.

MP = 156 - 158°C;

 $^1\text{H NMR}$ (CDCl_3 , 300 MHz) δ 7.70-7.26 (m, 9H), 5.28-5.12 (m, 1H), 4.57 (s, 1H), 4.35-4.10 (m, 1H). $[\alpha]_D = -71.2^\circ$ ($c=0.66$, MeOH)

Step F: Preparation of (+)-3,5-Dichloro-N-[3R-2,3-dihydro-2-oxo-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]benzamide:

To a stirring solution of (+)-3R-3-amino-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4] diazepine (5.6 g, 16.8 mmol) in DMF (50 mL) was added 1-(3-Dimethylaminopropyl-3-ethylcarbodi-imide hydrochloride (4.44 g, 23.0 mmol), and 1-hydroxybenztriazole hydrate (3.11 g, 23.0 mmol) and 3,5-Dichlorobenzoic acid (3.21 g, 16.8 mmol). This was stirred at ambient temperature for 2 hours. The reaction was diluted with 500 mL satd. NaHCO_3 and extracted with 2 x 300 mL ethyl acetate. The combined organics were washed with 10% KHSO_4 (200 mL), brine (200 mL), dried over Na_2SO_4 , and evaporated to a white foam. This was chromatographed over a 75 x 200 mm silica column eluting with 20% ethyl acetate:hexane. The pure fractions were collected and evaporated under reduced pressure to give 8.5 g of a white foam which was crystallized from 15% ethyl acetate:hexane to give 5.3 g of a white powder. mp=140-143°C, $[\alpha]_D = +47.9^\circ$; $^1\text{H NMR}$ (CDCl_3 , 300 MHz) δ 7.85-7.75 (m, 2 H), 7.70-7.20 (m, 9 H), 5.78 (d, $J=8.1$ Hz, 1 H), 5.30-5.15 (m, 1H), 4.30-4.15 (m, 1H)

Analysis Calcd. for $\text{C}_{24}\text{H}_{16}\text{Cl}_2\text{F}_3\text{N}_3\text{O}_2$:

C, 56.93; H, 3.19; N, 8.30;

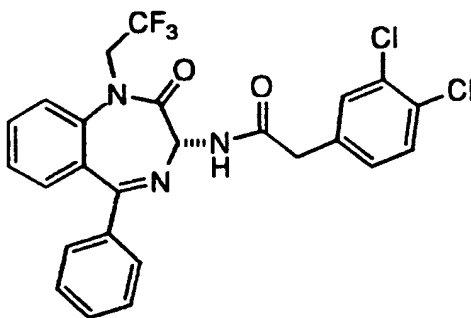
Found: C, 56.81; H, 3.17; N, 8.17.

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The following examples were prepared by a procedure substantially as described for Example 1, Step F.

EXAMPLE 113

(-)-2-(3,4-Dichlorophenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzof[e][1,4]diazepin-3-yl]acetamide.



mp=219-221°C; $[\alpha]_D = -10.8^\circ$;

^1H NMR (CDCl_3 , 300 MHz) δ 7.65-7.15 (m, 12H), 5.78 (d, $J=8.1$ Hz, 1H), 5.25-5.10 (m, 1H), 4.25-4.05 (m, 1H), 3.56 (s, 2H);

Analysis Calcd. for $\text{C}_{25}\text{H}_{18}\text{Cl}_2\text{F}_3\text{N}_3\text{O}_2 \cdot 0.85 \text{H}_2\text{O}$:

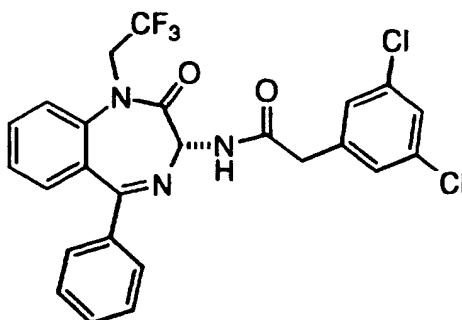
C, 56.06; H, 3.71; N, 7.84.

Found: C, 56.03; H, 3.53; N, 7.82.

EXAMPLE 114

(-)-2-(3,5-Dichlorophenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzof[e][1,4]diazepin-3-yl]acetamide

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mp=93-100°C, $[\alpha]_D = -5.7^\circ$;

$^1\text{H NMR}$ (CDCl_3 , 300 MHz) δ 7.65-7.15 (m, 12H), 5.78 (d, $J=8.1$ Hz, 1H), 5.25-5.10 (m, 1H), 4.25-4.05 (m, 1H), 3.65 (s, 2H);

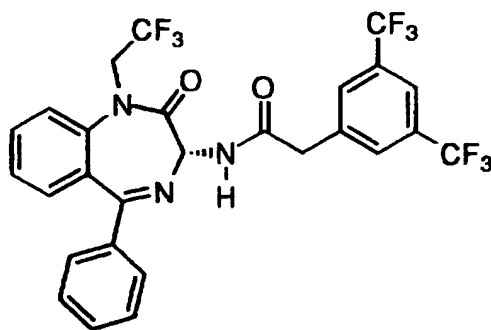
Analysis Calcd. for $\text{C}_{25}\text{H}_{18}\text{Cl}_2\text{F}_3\text{N}_3\text{O}_2$:

C, 57.71; H, 3.49; N, 8.08;

Found : C, 57.41; H, 3.48; N, 8.12.

EXAMPLE 115

(-)-2-[3,5-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



m.p. foam °C, $[\alpha]_D = -9.7^\circ$ ($c=0.59$, MeOH).

Anal. Calcd. for $\text{C}_{27}\text{H}_{18}\text{F}_9\text{N}_3\text{O}_2 \cdot 0.75 \text{H}_2\text{O}$:

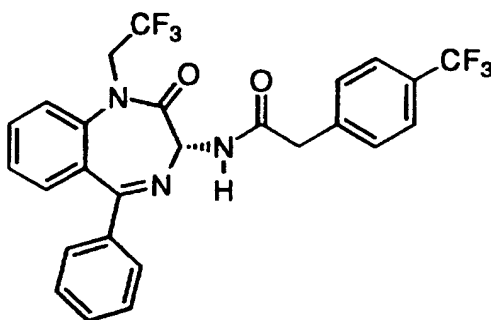
C, 53.96; H, 3.27; N, 6.99.

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Found: C, 53.96; H, 3.1; N, 6.98%.

EXAMPLE 116

(-)-2-(4-Trifluoromethylphenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



m.p. 253-255 °C, $[\alpha]_D = -9.2^\circ$ ($c=0.25$, MeOH).

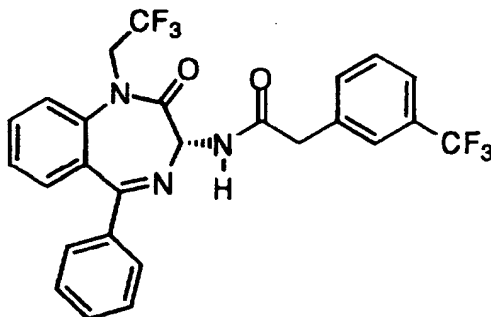
Anal. Calcd. for $C_{26}H_{19}F_6N_3O_2 \cdot 0.05$ ethyl ether 0.55 H₂O:

C, 59.03; H, 3.9; N, 7.88.

Found: C, 59.05; H, 3.82; N, 7.78%.

EXAMPLE 117

2-(3-Trifluoromethylphenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



m.p. 172-173 °C, $[\alpha]_D = +5.9^\circ$ ($c=0.56$, CHCl₃).

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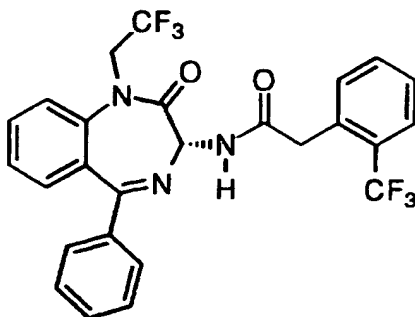
Anal. Calcd. for $C_{26}H_{19}F_6N_3O_2 \cdot 0.60 H_2O$:

C, 58.89; H, 3.84; N, 7.92.

Found: C, 58.92; H, 3.71; N, 7.98%.

EXAMPLE 118

(+)-2-(2-Trifluoromethylphenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide

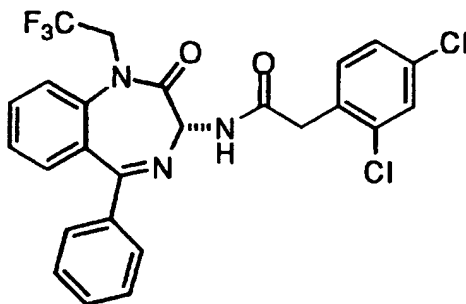
m.p. 170-171 °C, $[\alpha]_D = +9.0^\circ$ (c=0.48, $CHCl_3$).Anal. Calcd. for $C_{26}H_{19}F_6N_3O_2 \cdot 0.25 H_2O$:

C, 59.6; H, 3.75; N, 8.02.

Found: C, 59.64; H, 3.68; N, 7.97%.

EXAMPLE 119

(-)-2-(2,4-Dichlorophenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



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m.p. 143-145 °C, $[\alpha]_D = -22.6^\circ$ ($c=0.73$; MeOH).

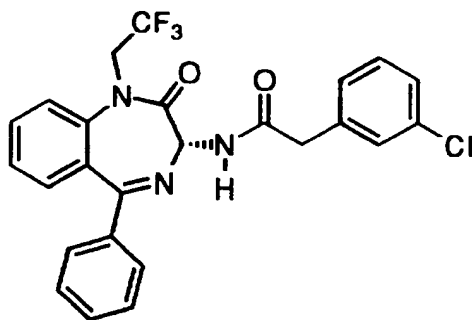
Anal. Calcd. for $C_{25}H_{18}N_3O_2Cl_2F_3$:

C, 57.71; H, 3.49; N, 8.08.

Found: C, 57.75; H, 3.52; N, 8.09%.

EXAMPLE 120

(-)-2-(3-Chlorophenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



m.p. 188-189 °C, $[\alpha]_D = -5.4^\circ$ ($c=1.03$, MeOH).

Anal. Calcd. for $C_{25}H_{19}ClF_3N_3O_2 \cdot 0.10$ ethyl ether:

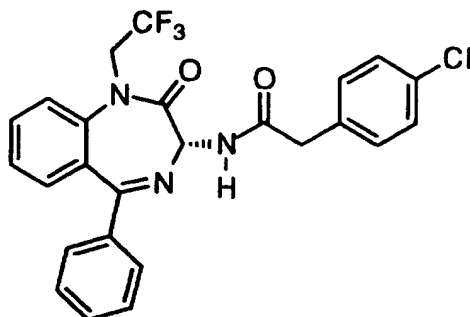
C, 61.84; H, 4.09; N, 8.52.

Found: C, 61.84; H, 4.05; N, 8.5%.

EXAMPLE 121

(-)-2-(4-Chlorophenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide

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m.p. 246-247 °C, $[\alpha]_D = -10.1^\circ$ ($c=0.45$, MeOH).

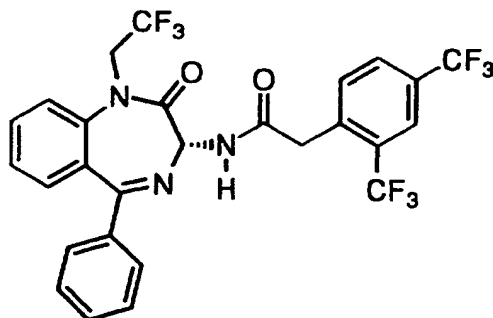
Anal. Calcd. for $C_{25}H_{19}ClF_3N_3O_2 \cdot 0.20 H_2O \cdot 0.15$ ethyl ether:

C, 61.42; H, 4.21; N, 8.39.

Found: C, 61.46; H, 4.15; N, 8.39%.

Example 122

(-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



Step A. 2,4-Bis(trifluoromethyl)benzonitrile

To a stirring biphasic mixture of 100mL ethanol and 250 mL of phosphate buffer (1g of $NaH_2PO_4 \cdot H_2O$ per 5 mL H_2O adjusted to pH=7.0 with 50% NaOH) and NaCN (81.3mmol, 4.0g) heated to 60°C was added 2,4-bis(trifluoromethyl) benzyl bromide (32.5mmol, 10g) in 50mL EtOH dropwise over 30min. The reaction was heated at 60°C

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for 24h. The reaction was then evaporated under reduced pressure. The remaining aqueous was extracted with 2x150mL EtOAc. The organic layers were combined, dried with brine and Na₂SO₄. The organic phase was evaporated under reduced pressure and the residue chromatographed over silica eluting with 10% EtOAc:Hexanes. The pure fractions were collected and evaporated to give 7.0g of a pale yellow oil, 85.1% NMR ¹H (CDCl₃) δ 8.0-7.85 (m,3H), 4.03 (s,2H)

Step B. 2,4-Bis(trifluoromethyl)phenyl acetic acid

2,4-Bis(trifluoromethyl)benzonitrile (41.5mmol,10.51g) was taken up in 100mL acetic acid, 50mL conc. H₂SO₄, and 20mL water. This was heated to 120°C for 3h. The reaction was then diluted with 1L ice water, and extracted with 2x300mL ethyl acetate. The combined organics were washed with 2x200mL water, dried with brine and Na₂SO₄, and evaporated under reduced pressure. The residue was taken up in a minimum of diethyl ether and crystallized by adding sufficient hexane to precipitate the product. The solid was collected to give 7.74g of 2,4-bis(trifluoromethyl) phenyl acetic acid as white crystals, 68.5%. NMR ¹H (CDCl₃) δ 7.93 (s,1H), 7.80 (d, J=7.9Hz,1H), 7.55 (d, J=7.9Hz,1H), 3.94 (s,2H).

Step C. Preparation of (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide

To a stirring solution the 3R-3-Amino-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-2,3-dihydro-1H-benzo[e][1,4]diazepine (28.4 mmol, 9.47g) in DMF (100mL) was added 2,4-Bis(trifluoromethyl) phenyl acetic acid (28.4mmol,7.74g), 1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (42.6mmol,8.16g) and 1-Hydroxybenzotriazole hydrate (14.2mmol,1.92g). This was stirred for 1h at room temperature. The reaction was then diluted with 750mL of 10% KHSO₄ and extracted with ethyl acetate (2x300mL). The organic layers

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were combined and washed with saturated sodium hydrogen carbonate (1x600mL). The organics were then dried with brine, and sodium sulfate and evaporated under reduced pressure. The residue was chromatographed over silica eluting with 20% EtOAc:Hexane. Pure fractions were collected and evaporated. The residue was taken up in 100 mL of warm 75% isopropanol:water. This was allowed to cool slowly and stirred overnight (16 hr) at room temperature. The suspension was cooled briefly to @5°C and filtered. The white solid was dried overnight at 60°C to give 10.5 g of material that melted at 132-134°C. X-Ray diffraction confirms crystallinity.

NMR ^1H (CDCl_3) δ 7.95-7.25 (m, 13H), 5.60 (d, $J=8.1\text{Hz}$, 1H), 5.30-5.10 (m, 1H), 4.25-4.06 (m, 1H), 3.96 (s, 2H)

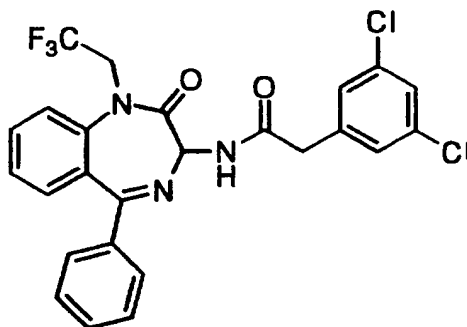
Anal. Calcd. for $\text{C}_{27}\text{H}_{18}\text{F}_9\text{N}_3\text{O}_2$:

C, 55.20; H, 3.09; N, 7.15.

Found: C, 55.03; H, 3.14; N, 7.10%.

EXAMPLE 123

(±)-2-(3,5-Dichlorophenyl)-N-[2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl] acetamide



m.p. 219-220 °C. racemic

Anal. Calcd. for $\text{C}_{25}\text{H}_{18}\text{N}_3\text{O}_2\text{Cl}_2\text{F}_3$:

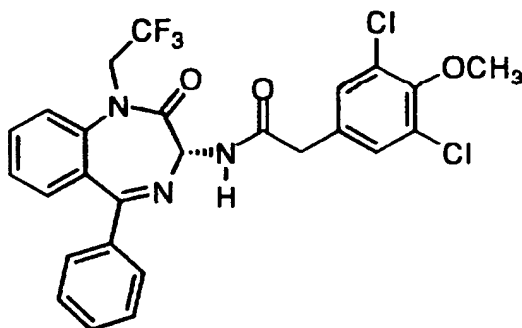
C, 57.71; H, 3.49; N, 8.08.

Found: C, 57.94; H, 3.48; N, 8.02%.

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EXAMPLE 124

2-(3,5-dichloro-4-methoxyphenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



m.p. 100-104 °C, $[\alpha]_D = -8.9^\circ$ (c=0.55, MeOH).

Anal. Calcd. for $C_{26}H_{20}Cl_2F_3N_3O_3$:

C, 56.74; H, 3.66; N, 7.63.

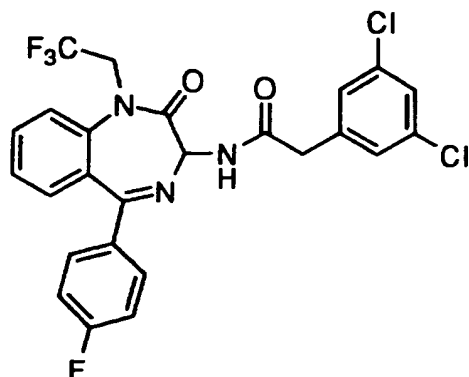
Found: C, 55.67; H, 3.47; N, 7.41%.

The following examples were prepared by procedures substantially as described in example 1 except substituting the appropriate fluoro substituted aminobenzophenone in step A.

EXAMPLE 125

(+)-2-(3,5-Dichlorophenyl)-N-[2,3-dihydro-5-(4-fluorophenyl)-2-oxo-1-(2,2,2-trifluoroethyl)-1H-benzof[e][1,4]diazepin-3-yl]acetamide

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m.p. foam °C, $[\alpha]_D = +3.4^\circ$ ($c=0.55$; MeOH).

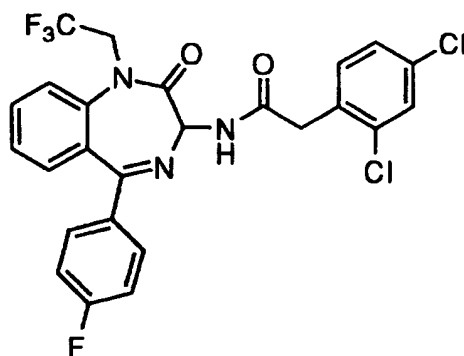
Anal. Calcd. for $C_{25}H_{17}N_3O_2Cl_2F_4$:

C, 55.78; H, 3.18; N, 7.81.

Found: C, 55.73; H, 3.25; N, 7.72%.

EXAMPLE 126

(-)-2-(2,4-Dichlorophenyl)-N-[2,3-dihydro-5-(4-fluorophenyl)-2-oxo-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



m.p. foam °C, $[\alpha]_D = -11^\circ$ ($c=0.68$; MeOH).

Anal. Calcd. for $C_{25}H_{17}N_3O_2F_4$:

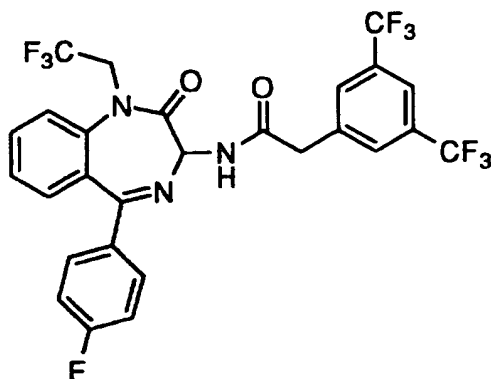
C, 55.78; H, 3.18; N, 7.81.

Found: C, 55.82; H, 3.41; N, 7.42%.

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EXAMPLE 127

(+)-2-(3,5-Bis(trifluoromethyl)phenyl)-N-[2,3-dihydro-5-(4-fluorophenyl)-2-oxo-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]-acetamide



m.p. foam °C, $[\alpha]_D = +2.8^\circ$ ($c=0.67$; MeOH).

Anal. Calcd. for $C_{27}H_{17}N_3O_2F_{10}$:

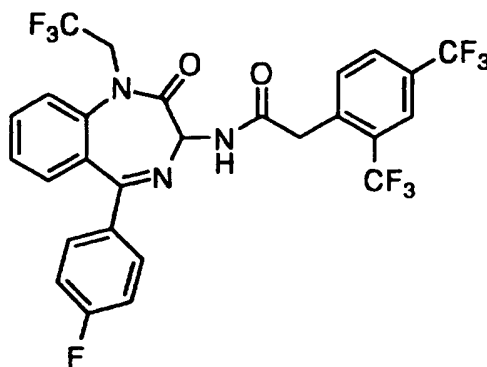
C, 53.56; H, 2.83; N, 6.94.

Found: C, 53.56; H, 2.93; N, 6.91%.

EXAMPLE 128

(-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[2,3-dihydro-5-(4-fluorophenyl)-2-oxo-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide

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$[\alpha]_D = -14^\circ$ ($c=0.63$; MeOH).

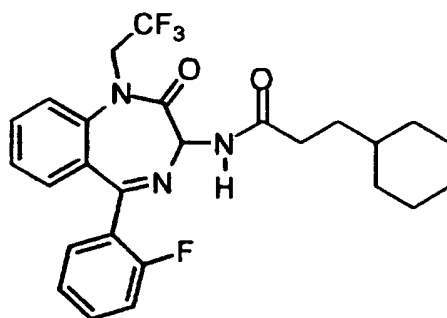
Anal. Calcd. for $C_{27}H_{17}N_3O_2F_{10}$:

C, 53.56; H, 2.83; N, 6.94.

Found: C, 53.3; H, 2.89; N, 7.05%.

EXAMPLE 129

3-Cyclohexyl-N-[2,3-dihydro-5-(2-fluorophenyl)-2-oxo-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]propionamide



m.p. 202-204 °C.

1H NMR δ ($CDCl_3$) 7.72 (m, 8H), 5.65 (d, $J=8.3$ Hz, 1H), 5.35-5.08 (m, 1H), 4.32-4.15 (m, 1H), 2.37 (t, $J=7.8$ Hz, 2H), 1.80-1.55 (m, 7H),

1.45. Anal. Calcd. for $C_{26}H_{27}F_4N_3O_2$:

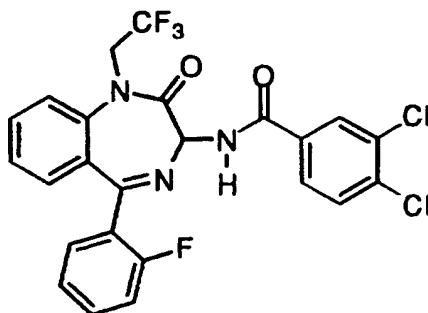
C, 63.8; H, 5.56; N, 8.58.

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Found: C, 63.82; H, 5.54; N, 8.56%.

EXAMPLE 130

3,4-Dichloro-N-[2,3-dihydro-5-(2-fluorophenyl)-2-oxo-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]benzamide



m.p. 168-170 °C.

¹H NMR δ (CDCl₃) 8.03 (d, J=2.0, 1H), 7.86 (d, J=7.8 Hz, 1H), 7.78-7.05 (m, 9H), 5.80 (d, J=7.8 Hz, 1H), 5.27-5.15 (m, 1H), 4.35-4.20 (m, 1H)

Anal. Calcd. for C₂₄H₁₅Cl₂F₄N₃O₂:

C, 54.98; H, 2.88; N, 8.01.

Found: C, 54.96; H, 2.89; N, 8.12%.

EXAMPLE 131

Antiarrhythmic efficacy of the combined administration of low dose I_{Ks} blocker (-)-2-[2,4-bis(trifluoromethyl)phenyl]-n-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide and the beta-adrenergic receptor antagonist timolol in anesthetized dogs with previous anterior myocardial infarction was studied. This study was conducted to test the hypothesis the concomitant administration of low dose I_{Ks} blocker (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide and

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low dose beta-adrenergic receptor antagonist timolol, which when administered individually at low dose afford no significant anti-arrhythmic protection, would in combination provide significant protection against the development of malignant ventricular arrhythmias in a canine model of previous myocardial infarction. In the absence of protective intervention, this postinfarction canine preparation displays a very high incidence (80%) of malignant ventricular tachycardia degenerating into ventricular fibrillation and death in response to the development of ischemia at a site remote from previous myocardial infarction (Patterson et al., Am. J. Cardiol. 50: 1414-1423, 1982; Wilber et al., Am. Heart J. 109: 8-18, 1985; Lynch et al., J Pharmacol Exp Therap 277: 671-678, 1996).

METHODS

Surgical Preparation.

Male or female purpose-bred mongrel dogs (7-12 kg) were preanesthetized with sodium thiamylal (5.0 mg/kg i.v.) and general anesthesia was induced with isoflurane. A left thoracotomy was performed in the 4th intercostal space, the pericardium incised, and the heart suspended in a pericardial cradle. Anterior myocardial infarction was produced by a two hour occlusion of the left anterior descending coronary artery followed by reperfusion. Surgical incisions were closed, and the animals were allowed to recover.

Electrophysiologic Testing, Programmed Ventricular Stimulation (PVS) and Acute Posterolateral Myocardial Ischemia

Purpose-bred male or female mongrel dogs at 5-20 days after anterior myocardial infarction were anesthetized with alpha chloralose (80.0-100.0 mg/kg i.v.). The animals were ventilated by means of a cuffed endotracheal tube and a volume-cycled respirator (Model 613, Harvard Apparatus, S. Natick, MA). Systemic arterial pressure was monitored via the cannulated left common carotid artery (Model P23XL pressure transducer, Gould Inc, Cleveland, OH), and the

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right femoral vein was isolated and cannulated for drug administration. The heart was re-exposed via a left thoracotomy, and was suspended in a pericardial cradle. A platinum epicardial bipolar electrode (3 mm electrode post separation) was sutured to the surface of the left atrial appendage for atrial pacing, and a stainless steel bipolar plunge electrode (5 mm length, 3 mm electrode post separation) was inserted into the interventricular septum near the right ventricular outflow tract (RVOT) adjacent to the site of left anterior descending coronary artery occlusion for the introduction of ventricular extrastimuli during programmed ventricular stimulation (PVS). One acrylic button per zone containing fixed stainless steel bipolar plunge electrodes (see below for specifications) was sutured into the infarcted anterior region of the left ventricle distal to the site of coronary artery occlusion and within the area of myocardial scarring as ascertained visually and by palpation (IZ, infarct zone) and into the non-infarcted posterolateral region of the left ventricle (NZ, non-infarct zone) for the measurement of ventricular electrophysiologic parameters (excitation thresholds and refractory periods). Lead II electrocardiogram was monitored continuously.

After stabilization of the preparation and the measurement of baseline electrocardiographic and cardiac electrophysiologic parameters, PVS consisting of the introduction of 1-3 ventricular extrastimuli during sinus rhythm and atrial pacing was performed at the RVOT site. If ventricular extrastimuli could not be inserted at the RVOT site, programmed pacing was attempted at the IZ site. Ventricular extrastimuli were introduced at 2x diastolic threshold voltage or, if extrastimuli were not introduced adequately, at 4x diastolic threshold voltage. Responses to baseline programmed ventricular stimulation were categorized as: 1) non-inducible (NI): less than 5 nonstimulated ventricular complexes; 2) nonsustained ventricular tachycardia (NSVT): 5 or more nonstimulated ventricular complexes terminating spontaneously with a duration less than 15 seconds; 3) unimorphic sustained ventricular tachycardia (UVT): unimorphic ventricular tachycardia with a duration exceeding 15 seconds; 4) polymorphic sustained ventricular tachycardia (PVT):

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polymorphic ventricular tachycardia with a duration exceeding 15 seconds, and 5) ventricular tachycardia degenerating into ventricular fibrillation (VT/VF): an unstable ventricular tachycardia degenerating spontaneously into ventricular fibrillation. In this study, programmed pacing was continued until the induction of either sustained VT or VT/VF, or until the end of the pacing protocol with either NSVT or no response (NI) occurring. In many preparations ultimately responding to programmed stimulation with either SVT or VT/VF, reproducible NSVTs were initiated early in the pacing protocol. Only postinfarction preparations responding to baseline PVS with inducible sustained ventricular tachyarrhythmias (UVT, PVT or VT/VF) were entered into the present study. Baseline response to PVS with inducible ventricular tachyarrhythmias has been found to correlate strongly with the development malignant ventricular tachycardia, fibrillation and arrhythmic death in response to a subsequent episode of myocardial ischemia at a site remote from previous infarction in this animal model (Wilber et al., Am. Heart J. 109: 8-18, 1985).

Following the completion of baseline and post treatment electrophysiologic and PVS testing, the tip of a silver wire electrode was inserted through the wall and into the lumen of the proximal left circumflex (LCX) coronary artery. An anodal current of 200 μ A was applied to the intimal surface of the coronary artery via this electrode, producing intimal injury, thrombus formation and ultimately acute myocardial ischemia in the left circumflex coronary artery distribution. The onset of acute posterolateral myocardial ischemia was noted electrocardiographically. Upon the development of lethal ischemic arrhythmias or at 3 hours after the onset of acute posterolateral myocardial ischemia, the hearts were excised and wet thrombus mass in the left circumflex coronary artery determined. Anterior myocardial infarct size was determined by cutting the heart into 1 cm thick transverse sections which then were incubated in 0.4% triphenyltetrazolium chloride solution. Reaction with triphenyltetrazolium forms a red precipitate in viable tissue, whereas infarcted tissue remains pale. Infarct size was

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quantitated gravimetrically and was expressed as a percentage of total left ventricle.

The responses (i.e. occurrence of vs protection from the development of lethal ventricular arrhythmia) of five experimental groups to the development of acute posterolateral ischemia at a site remote from previous myocardial infarction were compared in this study: 1) the laboratory cohort of aqueous vehicle (saline or PEG/saline)-treated control animals (n=40); 2) a dedicated soy bean oil-based microemulsion vehicle (20.0% soy bean oil, 2.0% glycerin, 1.2% lecithin and 76.8% water)-treated control group (n=10); 3) low dose 0.0003 mg/kg i.v. I_{K_S} blocker (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide in soy bean oil-based microemulsion (n=10); 4) low dose 0.001 mg/kg i.v. beta-adrenergic receptor blocking agent timolol in 5% dextrose in saline vehicle (n=6); and 5) the combination of low dose 0.0003 mg/kg i.v. I_{K_S} blocker (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide with low dose 0.001 mg/kg i.v. beta-adrenergic receptor blocking agent timolol (n=10). In the individual and combination I_{K_S} blocker (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide and beta-adrenergic receptor blocking agent timolol treatment groups, treatments were administered i.v. 15 min prior to the conduct of post treatment PVS and production of ischemia at a site remote from previous infarction.

Statistics.

Data are expressed as mean \pm S.E.M. Comparisons among treatment groups were made using a single factor ANOVA or a Fisher's exact test, when appropriate. P 0.05 was the criterion for statistical significance.

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RESULTS

All animals in all treatment groups responded to baseline PVS with inducible ventricular tachyarrhythmias as an entry criterion. The incidences of ischemic lethal ventricular arrhythmias (ventricular fibrillation; VF) occurring in the laboratory control cohort, dedicated microemulsion vehicle control group, low dose I_{Ks} blocker (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide, low dose beta-adrenergic receptor blocking agent timolol, and combined low dose I_{Ks} blocker (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide with beta-adrenergic receptor blocking agent timolol groups are summarized below.

Treatment Group	Incidents of Lethal Ischemic Ventricular Arrhythmic (VF)	P Value Compared to Laboratory Vehicle Control Cohort	P Value Compared to Microemulsion Vehicle Control Group
Laboratory Vehicle Control Cohort	34/40 (85%)	--	NS
Microemulsion Vehicle Control Group	8/10 (80%)	NS	--
Low Dose I_{Ks} Blocker 0.0003 mg/kg i.v.	6/10 (60%)	NS	NS
Low Dose β Blocker 0.001 mg/kg i.v. Timolol	4/6 (67%)	NS	NS
Combined Low Dose I_{Ks} Blocker 0.0003 mg/kg i.v. + Low Dose β Blocker 0.001 mg/kg i.v. Timolol	3/10 (30%)	0.001	0.03

NS = nonsignificant ($P > 0.05$).

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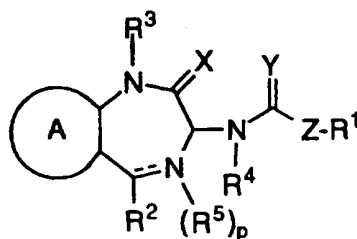
Neither low dose I_{K_S} blocker (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide nor low dose beta-adrenergic receptor antagonist timolol administered individually afforded significant protection against the development of malignant ischemic ventricular arrhythmias and death compared to either the laboratory vehicle control cohort or the dedicated microemulsion vehicle control group. In contrast, the combined administration of both low dose I_{K_S} blocker (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide and low dose beta-adrenergic receptor antagonist timolol provided significant protection against development of malignant ischemic ventricular tachyarrhythmia and arrhythmic death, which signifies a synergistic salutary antiarrhythmic interaction between block of I_{K_S} and beta-adrenergic receptor blockade.

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WHAT IS CLAIMED IS:

1. A method of treating arrhythmia which comprises the co-administration to a patient in need of such treatment of an effective amount of a beta-adrenergic receptor blocking agent and a selective I_{Ks} antagonist.

2. The method of Claim 1 wherein the selective I_{Ks} antagonist is a compound of structural formula:



I

individual diastereomers, enantiomers and mixtures thereof,
or a pharmaceutically acceptable salt thereof, wherein

A is

- 1) thieno,
- 2) pyrido, or
- 3) benzo either unsubstituted or substituted with $-NH_2$, $-NHSO_2(C_{1-3} \text{ alkyl})$, $C_{1-3} \text{ alkyl}$ or $C_{1-3} \text{ alkoxy}$;

X is

- 1) = O,
- 2) = S,
- 3) = N-NH₂,
- 4) = N-OH or
- 5) = H₂;

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Y is

- 1) = O,
- 2) = N-CN or
- 3) = H₂;

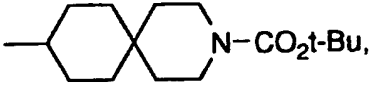
Z is

- 1) C₁-6 alkylene, either straight or branch chain and either unsubstituted or substituted with phenyl or spiropiperidine,
- 2) C₂-4 alkenylene, either straight or branch chain,
- 3) -(CH₂)_m-W-(CH₂)_n- wherein m and n are independently 0, 1, 2, 3 or 4 and W is -O-, -S- or -NH,
- 4) 4-(5-methylisoxazole-3-yl),
- 5) C₃-6 cycloalkylene, or
- 6) single bond;

p is 0 or 1;

R¹ is

- 1) phenyl, either unsubstituted or substituted with one or two substituents selected from
 - a) -NO₂,
 - b) -Cl, Br, F, or I,
 - c) -CF₃,
 - d) -C₁-3 alkyl,
 - e) -C₁-3 alkoxy,
 - f) -CN,
 - g) -methylenedioxy,
- 2) C₅-7 cycloalkyl,
- 3)


- 4) mono- or bicyclic heterocyclyl of 5 to 10 members one or two of which are sulfur, nitrogen or oxygen, the remaining

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being carbon, such as 2-thienyl, 2-furanyl, 2-indolyl, 2-quinoxoliny, or 2-(2,3-dihydro benzofuranyl)

- 5) methyl, or
- 6) indan-5-yl;

R² is

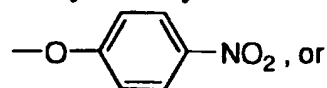
- 1) phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,
- 2) C₁₋₄ alkyl, either straight or branched chain and either unsubstituted or substituted with C₁₋₃ alkoxy or C₁₋₃ alkoxy-C₁₋₃ alkoxy,
- 3) C₅₋₇ cycloalkyl,
- 4) 2- or 3-furyl,
- 5) 1-methylpiperidin-2-yl, or
- 6) if R² is phenyl, the 2-position of the phenyl can be joined to the 4-position nitrogen of the diazepine ring through a carbonyl group and the double bond between the 4-nitrogen and the 5-carbon becomes a single bond;

R³ is

- 1) hydrogen or
- 2) C₁₋₆ alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃, or
- 3) -CF₃;

R⁴ is

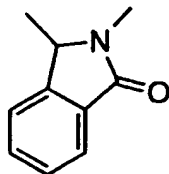
- 1) hydrogen,
- 2) C₁₋₆ alkyl, the chain of carbon atoms of which can be interrupted by one or two non-adjacent oxygen atoms and which is either unsubstituted or substituted with C₁₋₃ alkoxy, carbonyl, -OH or



- 3) tetrazol-5-yl; and

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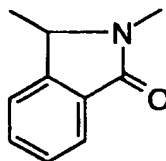
R^5 is hydrogen or oxygen or is joined to R^2 to form the partial structure:



;

and the bond represented by ---- is:

- 1) a double bond when p is zero or when p is 1 and R^5 is oxygen, or
- 2) a single bond when R^5 is hydrogen or R^5 is joined to R^2 to form the partial structure:



3. The selective I_{K_S} compound of Claim 2 wherein

A is benzo either unsubstituted or substituted with $-NH_2$, $-NHSO_2(C_1-3 \text{ alkyl})$, $C_1-3 \text{ alkyl}$ or $C_1-3 \text{ alkoxy}$;

X and Y are O,

Z is

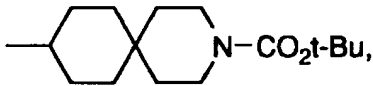
- 1) C_1-6 alkylene, either straight or branch chain and either unsubstituted or substituted with phenyl or spiropiperidine,
- 2) C_2-4 alkenylene, either straight or branch chain,
- 3) $-(CH_2)_m-W-(CH_2)_n-$ wherein m and n are independently 0, 1, 2, 3 or 4 and W is $-O-$, $-S-$ or $-NH$,
- 4) 4-(5-methylisoxazole-3-yl),
- 5) C_3-6 cycloalkylene, or

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- 6) single bond;

R¹ is

- 1) phenyl, either unsubstituted or substituted with one or two substituents selected from
 - a) -NO₂,
 - b) -Cl, Br, F, or I,
 - c) -CF₃,
 - d) -C₁₋₃ alkyl,
 - e) -C₁₋₃ alkoxy,
 - f) -CN,
 - g) -methylenedioxy,
- 2) C₅₋₇ cycloalkyl,
- 3)


*C1CCC2CCCCN2C1C(=O)OCC
- 4) mono- or bicyclic heterocyclyl of 5 to 10 members one or two of which are sulfur, nitrogen or oxygen, the remaining being carbon, such as 2-thienyl, 2-furanyl, 2-indolyl, 2-quinoxoliny, or 2-(2,3-dihydro benzofuranyl)
- 5) methyl, or
- 6) indan-5-yl;

R² is

- 1) phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,
- 2) C₁₋₄ alkyl, either straight or branched chain and either unsubstituted or substituted with C₁₋₃ alkoxy or C₁₋₃ alkoxy-C₁₋₃ alkoxy,
- 3) C₅₋₇ cycloalkyl,
- 4) 2- or 3-furyl,
- 5) 1-methylpiperidin-2-yl, or

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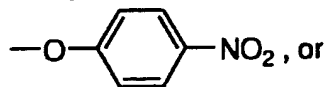
- 6) if R^2 is phenyl, the 2-position of the phenyl can be joined to the 4-position nitrogen of the diazepine ring through a carbonyl group and the double bond between the 4-nitrogen and the 5-carbon becomes a single bond;

 R^3 is

- 1) hydrogen or
- 2) C₁-6 alkyl, either straight or branched chain and either unsubstituted or substituted with $-N(CH_3)_2$, $-OH$, $-CF_3$, or
- 3) $-CF_3$;

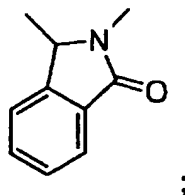
 R^4 is

- 1) hydrogen,
- 2) C₁-6 alkyl, the chain of carbon atoms of which can be interrupted by one or two non-adjacent oxygen atoms and which is either unsubstituted or substituted with C₁-3 alkoxy carbonyl, $-OH$ or



- 3) tetrazol-5-yl; and

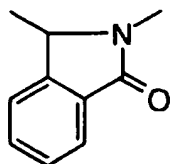
R^5 is hydrogen or oxygen or is joined to R^2 to form the partial structure:



and the bond represented by ---- is:

- 1) a double bond when p is zero or when p is 1 and R^5 is oxygen, or
- 2) a single bond when R^5 is hydrogen or R^5 is joined to R^2 to form the partial structure:

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including individual diastereomers, enantiomers and mixtures thereof, or a pharmaceutically acceptable salt.

4. The selective I_{K_S} compound of Claim 3 wherein

A is benzo either unsubstituted or substituted with $-NH_2$, $-NHSO_2(C_{1-3} \text{ alkyl})$, $C_{1-3} \text{ alkyl}$ or $C_{1-3} \text{ alkoxy}$;

X and Y are O,

Z is

- 1) C_{1-6} alkylane, either straight or branch chain and either unsubstituted or substituted with phenyl or spiropiperidine,
- 2) C_{2-4} alkenylene, either straight or branch chain,
- 3) $-(CH_2)_m-W-(CH_2)_n-$ wherein m and n are independently 0, 1, 2, 3 or 4 and W is $-O-$, $-S-$ or $-NH$,
- 4) 4-(5-methylisoxazole-3-yl),
- 5) C_{3-6} cycloalkylene, or
- 6) single bond;

R^1 is phenyl, either unsubstituted or substituted with one or two substituents selected from

- a) $-NO_2$,
- b) $-Cl$, Br , F , or I ,
- c) $-CF_3$,
- d) $-C_{1-3} \text{ alkyl}$,
- e) $-C_{1-3} \text{ alkoxy}$,
- f) $-CN$,
- g) $-methylenedioxy$,

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R² is phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,

R³ is -CF₃ or C₁₋₆ alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃;

R⁴ and R⁵ are hydrogen;

including individual diastereomers, enantiomers and mixtures thereof, or a pharmaceutically acceptable salt.

5. The selective I_{Ks} compound of Claim 4 wherein A is unsubstituted benzo;

R¹ is phenyl, either unsubstituted or substituted with one or two substituents selected from

- a) -NO₂,
- b) -Cl, Br, F, or I,
- c) -CF₃,
- d) -C₁₋₃ alkyl,
- e) -C₁₋₃ alkoxy,
- f) -CN,
- g) -methylenedioxy,

R² is phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,

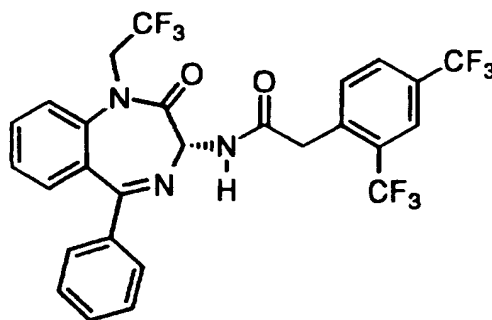
R³ is -CF₃ or C₁₋₆ alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃;

R⁴ and R⁵ are hydrogen;

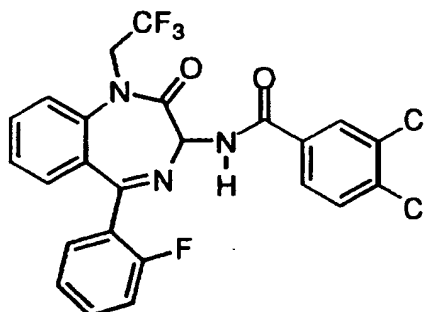
including individual diastereomers, enantiomers and mixtures thereof, or a pharmaceutically acceptable salt.

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6. The selective I_{K_S} compound of Claim 5 which is (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide

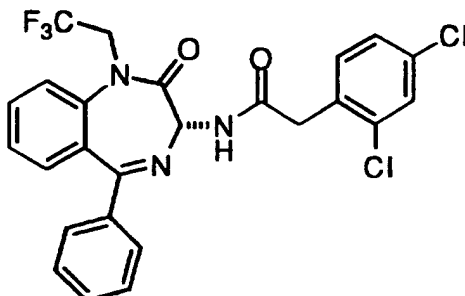


7. The selective I_{K_S} compound of Claim 5 which is 3,4-Dichloro-N-[2,3-dihydro-5-(2-fluorophenyl)-2-oxo-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]benzamide



8. The selective I_{K_S} compound of Claim 5 which is (-)-2-(2,4-Dichlorophenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide

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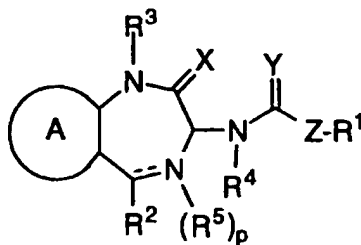


8. The beta-adrenergic receptor blocking agent of Claim 1 selected from the group consisting of timolol, sotalol, esmolol, catechol, propranolol, betaxolol, penbutolol, metoprolol, acebutolol, atenolol, metoprolol, pindolol, or bisoprolol.

9. A pharmaceutical formulation comprising a pharmaceutically acceptable carrier and a therapeutically effective amount of a beta-adrenergic receptor blocking agent and a selective I_{K_S} antagonist.

10. The beta-adrenergic receptor blocking agent of Claim 9 selected from the group consisting of timolol, sotalol, esmolol, catechol, propranolol, betaxolol, penbutolol, metoprolol, acebutolol, atenolol, metoprolol, pindolol, or bisoprolol.

11. The selective I_{K_S} antagonist of Claim 9 which is a compound of structural formula:



I

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individual diastereomers, enantiomers and mixtures thereof,
or a pharmaceutically acceptable salt thereof, wherein

A is

- 1) thieno,
- 2) pyrido, or
- 3) benzo either unsubstituted or substituted with -NH₂,
-NHSO₂(C₁₋₃ alkyl), C₁₋₃ alkyl or C₁₋₃ alkoxy;

X is

- 1) = O,
- 2) = S,
- 3) = N-NH₂,
- 4) = N-OH or
- 5) = H₂;

Y is

- 1) = O,
- 2) = N-CN or
- 3) = H₂;

Z is

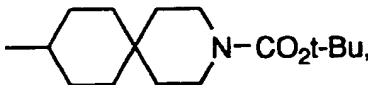
- 1) C₁₋₆ alkylene, either straight or branch chain and either unsubstituted or substituted with phenyl or spiropiperidine,
- 2) C₂₋₄ alkenylene, either straight or branch chain,
- 3) -(CH₂)_m-W-(CH₂)_n- wherein m and n are independently 0, 1, 2, 3 or 4 and W is -O-, -S- or -NH,
- 4) 4-(5-methylisoxazole-3-yl),
- 5) C₃₋₆ cycloalkylene, or
- 6) single bond;

p is 0 or 1;

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R¹ is

- 1) phenyl, either unsubstituted or substituted with one or two substituents selected from
 - a) -NO₂,
 - b) -Cl, Br, F, or I,
 - c) -CF₃,
 - d) -C₁₋₃ alkyl,
 - e) -C₁₋₃ alkoxy,
 - f) -CN,
 - g) -methylenedioxy,
- 2) C₅₋₇ cycloalkyl,
- 3)


CC(C)(C)OC(=O)N1CC2CCCCC2CC1
- 4) mono- or bicyclic heterocyclyl of 5 to 10 members one or two of which are sulfur, nitrogen or oxygen, the remaining being carbon, such as 2-thienyl, 2-furanyl, 2-indolyl, 2-quinoxoliny, or 2-(2,3-dihydro benzofuranyl)
- 5) methyl, or
- 6) indan-5-yl;

R² is

- 1) phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,
- 2) C₁₋₄ alkyl, either straight or branched chain and either unsubstituted or substituted with C₁₋₃ alkoxy or C₁₋₃ alkoxy-C₁₋₃ alkoxy,
- 3) C₅₋₇ cycloalkyl,
- 4) 2- or 3-furyl,
- 5) 1-methylpiperidin-2-yl, or
- 6) if R² is phenyl, the 2-position of the phenyl can be joined to the 4-position nitrogen of the diazepine ring through a carbonyl group and the double bond between the 4-nitrogen and the 5-carbon becomes a single bond:

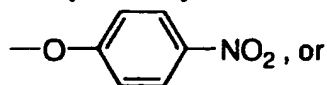
- 180 -

 R^3 is

- 1) hydrogen or
- 2) C₁-6 alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃, or
- 3) -CF₃;

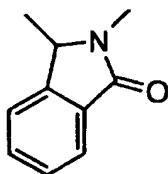
 R^4 is

- 1) hydrogen,
- 2) C₁-6 alkyl, the chain of carbon atoms of which can be interrupted by one or two non-adjacent oxygen atoms and which is either unsubstituted or substituted with C₁-3 alkoxycarbonyl, -OH or



- 3) tetrazol-5-yl; and

R^5 is hydrogen or oxygen or is joined to R^2 to form the partial structure:

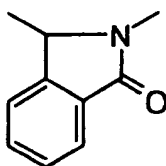


; and

the bond represented by ---- is:

- 1) a double bond when p is zero or when p is 1 and R^5 is oxygen, or
- 2) a single bond when R^5 is hydrogen or R^5 is joined to R^2 to form the partial structure:

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12. The selective I_{K_S} compound of Claim 11 wherein

A is benzo either unsubstituted or substituted with $-NH_2$, $-NHSO_2$ (C_{1-3} alkyl), C_{1-3} alkyl or C_{1-3} alkoxy;

X and Y are O,

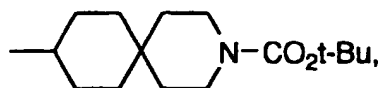
Z is

- 1) C_{1-6} alkylene, either straight or branch chain and either unsubstituted or substituted with phenyl or spiropiperidine,
- 2) C_{2-4} alkenylene, either straight or branch chain,
- 3) $-(CH_2)_m-W-(CH_2)_n-$ wherein m and n are independently 0, 1, 2, 3 or 4 and W is $-O-$, $-S-$ or $-NH$,
- 4) 4-(5-methylisoxazole-3-yl),
- 5) C_{3-6} cycloalkylene, or
- 6) single bond;

R^1 is

- 1) phenyl, either unsubstituted or substituted with one or two substituents selected from
 - a) $-NO_2$,
 - b) $-Cl$, Br , F , or I ,
 - c) $-CF_3$,
 - d) $-C_{1-3}$ alkyl,
 - e) $-C_{1-3}$ alkoxy,
 - f) $-CN$,
 - g) $-methylenedioxy$,
- 2) C_{5-7} cycloalkyl,
- 3)

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- 4) mono- or bicyclic heterocyclyl of 5 to 10 members one or two of which are sulfur, nitrogen or oxygen, the remaining being carbon, such as 2-thienyl, 2-furanyl, 2-indolyl, 2-quinoxolinyl, or 2-(2,3-dihydro benzofuranyl)
- 5) methyl, or
- 6) indan-5-yl;

R² is

- 1) phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,
- 2) C₁₋₄ alkyl, either straight or branched chain and either unsubstituted or substituted with C₁₋₃ alkoxy or C₁₋₃ alkoxy-C₁₋₃ alkoxy,
- 3) C₅₋₇ cycloalkyl,
- 4) 2- or 3-furyl,
- 5) 1-methylpiperidin-2-yl, or
- 6) if R² is phenyl, the 2-position of the phenyl can be joined to the 4-position nitrogen of the diazepine ring through a carbonyl group and the double bond between the 4-nitrogen and the 5-carbon becomes a single bond;

R³ is

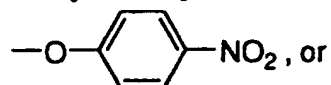
- 1) hydrogen or
- 2) C₁₋₆ alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃, or
- 3) -CF₃;

R⁴ is

- 1) hydrogen,
- 2) C₁₋₆ alkyl, the chain of carbon atoms of which can be interrupted by one or two non-adjacent oxygen atoms and

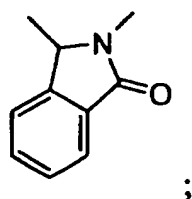
- 183 -

which is either unsubstituted or substituted with C₁-3
alkoxycarbonyl, -OH or



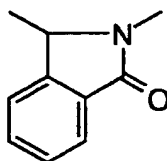
3) tetrazol-5-yl; and

R⁵ is hydrogen or oxygen or is joined to R² to form the partial structure:



and the bond represented by ---- is:

- 1) a double bond when p is zero or when p is 1 and R⁵ is oxygen, or
- 2) a single bond when R⁵ is hydrogen or R⁵ is joined to R² to form the partial structure:



including individual diastereomers, enantiomers and mixtures thereof,
or a pharmaceutically acceptable salt.

13. The selective I_K_S compound of Claim 12 wherein
A is benzo either unsubstituted or substituted with -NH₂, -NHSO₂(C₁-3
alkyl), C₁-3 alkyl or C₁-3 alkoxy;

X and Y are O,

Z is

- 184 -

- 1) C₁₋₆ alkylane, either straight or branch chain and either unsubstituted or substituted with phenyl or spiropiperidine,
- 2) C₂₋₄ alkenylene, either straight or branch chain,
- 3) -(CH₂)_m-W-(CH₂)_n- wherein m and n are independently 0, 1, 2, 3 or 4 and W is -O-, -S- or -NH,
- 4) 4-(5-methylisoxazole-3-yl),
- 5) C₃₋₆ cycloalkylene, or
- 6) single bond;

R¹ is phenyl, either unsubstituted or substituted with one or two substituents selected from

- a) -NO₂,
- b) -Cl, Br, F, or I,
- c) -CF₃,
- d) -C₁₋₃ alkyl,
- e) -C₁₋₃ alkoxy,
- f) -CN,
- g) -methylenedioxy,

R² is phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,

R³ is -CF₃ or C₁₋₆ alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃;

R⁴ and R⁵ are hydrogen;

including individual diastereomers, enantiomers and mixtures thereof, or a pharmaceutically acceptable salt.

14. The selective I_{K_S} compound of Claim 13 wherein A is unsubstituted benzo;

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R¹ is phenyl, either unsubstituted or substituted with one or two substituents selected from

- a) -NO₂,
- b) -Cl, Br, F, or I,
- c) -CF₃,
- d) -C₁₋₃ alkyl,
- e) -C₁₋₃ alkoxy,
- f) -CN,
- g) -methylenedioxy,

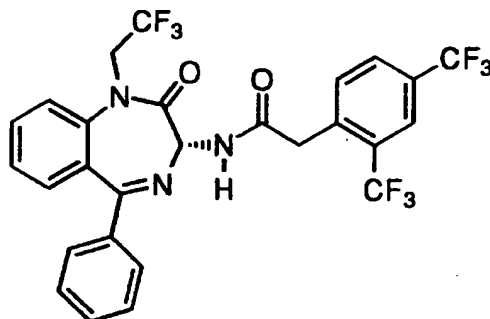
R² is phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,

R³ is -CF₃ or C₁₋₆ alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃;

R⁴ and R⁵ are hydrogen;

including individual diastereomers, enantiomers and mixtures thereof, or a pharmaceutically acceptable salt.

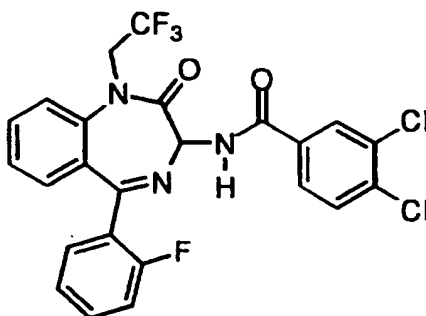
15. The selective I_{K_S} compound of Claim 14 which is (-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



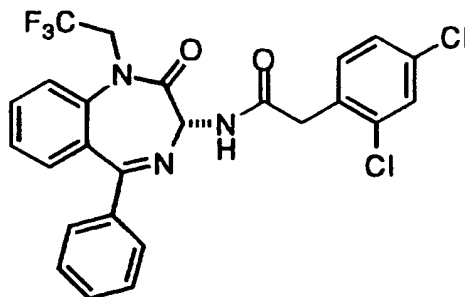
16. The selective I_{K_S} compound of Claim 14 which

- 186 -

is 3,4-Dichloro-N-[2,3-dihydro-5-(2-fluorophenyl)-2-oxo-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]benzamide



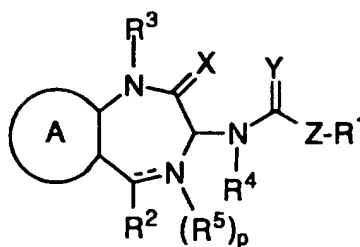
17. The selective I_{K_S} compound of Claim 14 which is (-)-2-(2,4-Dichlorophenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



18. A method of preventing arrhythmia which comprises the co-administration to a patient in need of such treatment of an effective amount of a beta-adrenergic receptor blocking agent and a selective I_{K_S} antagonist.

19. The method of Claim 18 wherein the selective I_{K_S} antagonist is a compound of structural formula:

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I

individual diastereomers, enantiomers and mixtures thereof,
or a pharmaceutically acceptable salt thereof, wherein

A is

- 1) thieno,
- 2) pyrido, or
- 3) benzo either unsubstituted or substituted with -NH₂,
-NHSO₂(C₁₋₃ alkyl), C₁₋₃ alkyl or C₁₋₃ alkoxy;

X is

- 1) = O,
- 2) = S,
- 3) = N-NH₂,
- 4) = N-OH or
- 5) = H₂;

Y is

- 1) = O,
- 2) = N-CN or
- 3) = H₂;

Z is

- 1) C₁₋₆ alkylene, either straight or branch chain and either
unsubstituted or substituted with phenyl or spiropiperidine,
- 2) C₂₋₄ alkenylene, either straight or branch chain,

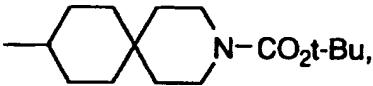
- 188 -

- 3) $-(CH_2)_m-W-(CH_2)_n-$ wherein m and n are independently 0, 1, 2, 3 or 4 and W is -O-, -S- or -NH,
- 4) 4-(5-methylisoxazole-3-yl),
- 5) C₃-6 cycloalkylene, or
- 6) single bond;

p is 0 or 1;

R¹ is

- 1) phenyl, either unsubstituted or substituted with one or two substituents selected from
 - a) -NO₂,
 - b) -Cl, Br, F, or I,
 - c) -CF₃,
 - d) -C₁₋₃ alkyl,
 - e) -C₁₋₃ alkoxy,
 - f) -CN,
 - g) -methylenedioxy,
- 2) C₅₋₇ cycloalkyl,
- 3)


- 4) mono- or bicyclic heterocyclyl of 5 to 10 members one or two of which are sulfur, nitrogen or oxygen, the remaining being carbon, such as 2-thienyl, 2-furanyl, 2-indolyl, 2-quinolyl, or 2-(2,3-dihydro benzofuranyl)
- 5) methyl, or
- 6) indan-5-yl;

R² is

- 1) phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,
- 2) C₁₋₄ alkyl, either straight or branched chain and either unsubstituted or substituted with C₁₋₃ alkoxy or

- 189 -

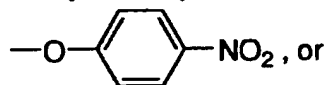
- C₁-3 alkoxy-C₁-3 alkoxy,
- 3) C₅-7 cycloalkyl,
 - 4) 2- or 3-furyl,
 - 5) 1-methylpiperidin-2-yl, or
 - 6) if R² is phenyl, the 2-position of the phenyl can be joined to the 4-position nitrogen of the diazepine ring through a carbonyl group and the double bond between the 4-nitrogen and the 5-carbon becomes a single bond;

R³ is

- 1) hydrogen or
- 2) C₁-6 alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃, or
- 3) -CF₃;

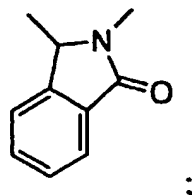
R⁴ is

- 1) hydrogen,
- 2) C₁-6 alkyl, the chain of carbon atoms of which can be interrupted by one or two non-adjacent oxygen atoms and which is either unsubstituted or substituted with C₁-3 alkoxy-carbonyl, -OH or



- 3) tetrazol-5-yl; and

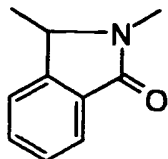
R⁵ is hydrogen or oxygen or is joined to R² to form the partial structure:



and the bond represented by ---- is:

- 190 -

- 1) a double bond when p is zero or when p is 1 and R⁵ is oxygen, or
- 2) a single bond when R⁵ is hydrogen or R⁵ is joined to R² to form the partial structure:



20. The selective I_{Ks} compound of Claim 19 wherein

A is benzo either unsubstituted or substituted with -NH₂, -NHSO₂(C₁-3 alkyl), C₁-3 alkyl or C₁-3 alkoxy;

X and Y are O,

Z is

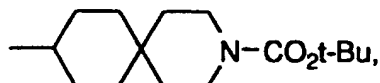
- 1) C₁-6 alkylene, either straight or branch chain and either unsubstituted or substituted with phenyl or spiropiperidine,
- 2) C₂-4 alkenylene, either straight or branch chain,
- 3) -(CH₂)_m-W-(CH₂)_n- wherein m and n are independently 0, 1, 2, 3 or 4 and W is -O-, -S- or -NH,
- 4) 4-(5-methylisoxazole-3-yl),
- 5) C₃-6 cycloalkylene, or
- 6) single bond;

R¹ is

- 1) phenyl, either unsubstituted or substituted with one or two substituents selected from
 - a) -NO₂,
 - b) -Cl, Br, F, or I,
 - c) -CF₃,

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- d) -C₁₋₃ alkyl,
- e) -C₁₋₃ alkoxy,
- f) -CN,
- g) -methylenedioxy,
- 2) C₅₋₇ cycloalkyl,
- 3)



- 4) mono- or bicyclic heterocyclyl of 5 to 10 members one or two of which are sulfur, nitrogen or oxygen, the remaining being carbon, such as 2-thienyl, 2-furanyl, 2-indolyl, 2-quinoxolinyl, or 2-(2,3-dihydro benzofuranyl)
- 5) methyl, or
- 6) indan-5-yl;

R² is

- 1) phenyl, either unsubstituted or substituted with C₁₋₃ alkoxy or 4,4-dimethyloxazolin-2-yl,
- 2) C₁₋₄ alkyl, either straight or branched chain and either unsubstituted or substituted with C₁₋₃ alkoxy or C₁₋₃ alkoxy-C₁₋₃ alkoxy,
- 3) C₅₋₇ cycloalkyl,
- 4) 2- or 3-furyl,
- 5) 1-methylpiperidin-2-yl, or
- 6) if R² is phenyl, the 2-position of the phenyl can be joined to the 4-position nitrogen of the diazepine ring through a carbonyl group and the double bond between the 4-nitrogen and the 5-carbon becomes a single bond;

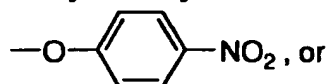
R³ is

- 1) hydrogen or
- 2) C₁₋₆ alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃, or
- 3) -CF₃;

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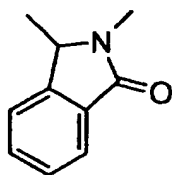
 R^4 is

- 1) hydrogen,
- 2) C_{1-6} alkyl, the chain of carbon atoms of which can be interrupted by one or two non-adjacent oxygen atoms and which is either unsubstituted or substituted with C_{1-3} alkoxy carbonyl, -OH or



- 3) tetrazol-5-yl; and

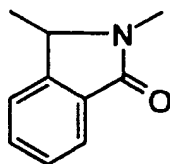
R^5 is hydrogen or oxygen or is joined to R^2 to form the partial structure:



; and

the bond represented by ---- is:

- 1) a double bond when p is zero or when p is 1 and R^5 is oxygen, or
- 2) a single bond when R^5 is hydrogen or R^5 is joined to R^2 to form the partial structure:



including individual diastereomers, enantiomers and mixtures thereof, or a pharmaceutically acceptable salt.

21. The selective IK_S compound of Claim 20 wherein

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A is benzo either unsubstituted or substituted with -NH₂, -NHSO₂ (C₁-3 alkyl), C₁-3 alkyl or C₁-3 alkoxy;

X and Y are O,

Z is

- 1) C₁-6 alkylane, either straight or branch chain and either unsubstituted or substituted with phenyl or spiropiperidine,
- 2) C₂-4 alkenylene, either straight or branch chain,
- 3) -(CH₂)_m-W-(CH₂)_n- wherein m and n are independently 0, 1, 2, 3 or 4 and W is -O-, -S- or -NH,
- 4) 4-(5-methylisoxazole-3-yl),
- 5) C₃-6 cycloalkylene, or
- 6) single bond;

R¹ is phenyl, either unsubstituted or substituted with one or two substituents selected from

- a) -NO₂,
- b) -Cl, Br, F, or I,
- c) -CF₃,
- d) -C₁-3 alkyl,
- e) -C₁-3 alkoxy,
- f) -CN,
- g) -methylenedioxy,

R² is phenyl, either unsubstituted or substituted with C₁-3 alkoxy or 4,4-dimethyloxazolin-2-yl,

R³ is -CF₃ or C₁-6 alkyl, either straight or branched chain and either unsubstituted or substituted with -N(CH₃)₂, -OH, -CF₃;

R⁴ and R⁵ are hydrogen;

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including individual diastereomers, enantiomers and mixtures thereof, or a pharmaceutically acceptable salt.

22. The selective I_{K_S} compound of Claim 21 wherein A is unsubstituted benzo;

R^1 is phenyl, either unsubstituted or substituted with one or two substituents selected from

- a) $-\text{NO}_2$,
- b) $-\text{Cl}$, Br , F , or I ,
- c) $-\text{CF}_3$,
- d) $-\text{C}_{1-3}$ alkyl,
- e) $-\text{C}_{1-3}$ alkoxy,
- f) $-\text{CN}$,
- g) $-\text{methylenedioxy}$,

R^2 is phenyl, either unsubstituted or substituted with C_{1-3} alkoxy or 4,4-dimethyloxazolin-2-yl,

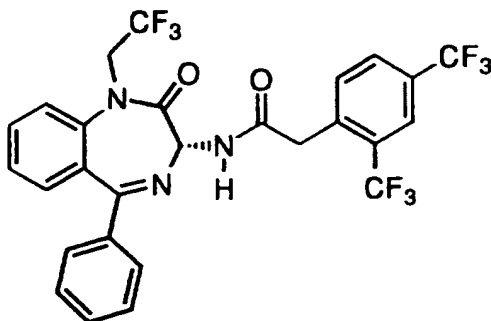
R^3 is $-\text{CF}_3$ or C_{1-6} alkyl, either straight or branched chain and either unsubstituted or substituted with $-\text{N}(\text{CH}_3)_2$, $-\text{OH}$, $-\text{CF}_3$;

R^4 and R^5 are hydrogen;

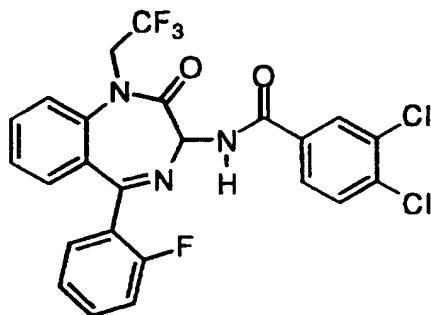
including individual diastereomers, enantiomers and mixtures thereof, or a pharmaceutically acceptable salt.

23. The selective I_{K_S} compound of Claim 22 which is
(-)-2-[2,4-Bis(trifluoromethyl)phenyl]-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide

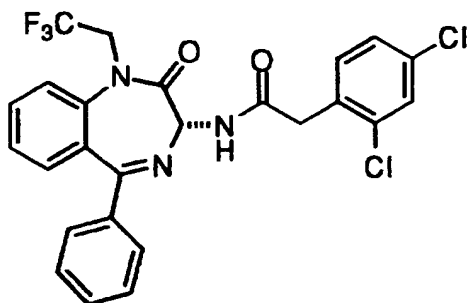
- 195 -



24. The selective I_{Ks} compound of Claim 22 which is 3,4-Dichloro-N-[2,3-dihydro-5-(2-fluorophenyl)-2-oxo-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]benzamide



25. The selective I_{Ks} compound of Claim 22 which is (-)-2-(2,4-Dichlorophenyl)-N-[3R-2,3-dihydro-2-oxo-5-phenyl-1-(2,2,2-trifluoroethyl)-1H-benzo[e][1,4]diazepin-3-yl]acetamide



26. The beta-adrenergic receptor blocking agent of Claim

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19 selected from the group consisting of timolol, sotalol, esmolol, cateolol, propranolol, betaxolol, penbutolol, metoprolol, acebutolol, atenolol, metoprolol, pindolol, or bisoprolol.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US97/11131

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : Please See Extra Sheet.

US CL : 540/494, 496, 501, 503, 504, 505, 509; 514/221

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 540/494, 496, 501, 503, 504, 505, 509; 514/221

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, CAS ONLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X, P	US 5,633,251 A (CLAREMON et al.) 27 May 1997, see entire document.	3-5, 11-14, 20-22
X	WO 96/05827 A1 (MERCK AND CO.) 29 February 1996, see entire document.	3-5, 11-14, 20-22
X	US 5,439,906 A (BOCK et al.) 08 August 1995, col. 7, line 12-17.	3-5, 11-14, 20-22
X	US 5,426,185 A (BALDWIN et al.) 20 June 1995, see entire document.	1-26
X	US 5,428,031 A (SANGUINETTI et al.) 27 June 1995, see entire document.	3-5, 11-14, 20-22
X	WO 95/14694 A1 (MERCK AND CO.) 01 June 1995, see entire document.	1-26



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

09 SEPTEMBER 1997

Date of mailing of the international search report

26 SEP 1997

Name and mailing address of the ISA/US
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Telephone No. (703) 308-1235

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/11131

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 96/05839 A1 (MERCK AND CO., INC.) 29 February 1996, see entire document.	1-26
X, P	US 5,556,969 A (CHAMBERS et al.) 17 September 1996, see entire document.	3-5, 11-14, 20-22
X	US 5,360,802 A (CHAMBERS et al.) 01 November 1994, see entire document.	3-5, 11-14, 20-22
X	US 5,324,726 A (BOCK et al.) 28 June 1994, col. 13, line 35 thru col. 14, line 38.	3-5, 11-14, 20-22
X	US 5,153,191 A (WOODRUFF) 06 October 1992, col. 1, lines 20-30 and claim 5.	3-5, 11-14, 20-22
X	US 4,994,258 A (BURNS et al.) 19 February 1991, see entire document.	3-5, 11-14, 20-22
X	US 4,628,084 A (BOCK et al.) 09 December 1986, see entire document.	3-5, 11-14, 20-22
X	US 4,970,207 A (SATO et al.) 13 November 1990, see entire document.	3-5, 11-14, 20-22
X	EP, 0, 167, 919 A2 (MERCK AND CO.) 15 January 1986, see entire document.	3-5, 11-14, 20-22
X	US 4,820,834 A (EVANS et al.) 11 April 1989, see entire document.	3-5, 11-14, 20-22
X, P	US 5,597,818 A (SANGUINETTI et al.) 28 January 1997, see entire document.	3-5, 11-14, 20-22
X, E	US 5,658,901 A (CLAREMON et al.) 19 August 1997, see entire document.	3-5, 11-14, 20-22
X, P	US 5,602,125 (CARLING et al.) 11 February 1997, see col. 11, lines 57-59.	11
X, P	US 5,595,990 A (BALDWIN et al.) 21 January 1997, see entire document.	3-5, 11-14, 20-22
X	US 5,521,175 A (CASTRO PINEIRO et al.) 28 May 1996, col. 17, lines 20-41.	3, 4, 11, 12, 20
X	US 5,451,582 A (CHAMBERS et al.) 19 September 1995, see entire document.	3, 4, 11, 12, 20

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/11131

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,439,905 A (NAKA et al.) 08 August 1995, see col. 5, lines 25-55.	11
X	US 5,220,017 A (BOCK et al.) 15 June 1993, see entire document.	3-5, 11-14, 20-22
X	US 4,992,437 A (NAKA et al.) 12 February 1991, see col. 6, lines 22-57.	11
X	WO 95/14694 A1 (MERCK AND CO.) 01 June 1995, see pages 10-13, 23, and 41.	3-5, 11-14, 20-22

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US97/11131

A. CLASSIFICATION OF SUBJECT MATTER:

IPC (6):

C07D 243/12, 243/18, 243/24, 255/04, 257/10, 259/00, 487/00, 491/00, 495/00; A61K 31/55